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Remarks:
The applicant has subsequently filed a sequence listing and declared, that it includes no new matter.

(54) Cellulose synthase gene

(57) mRNA was extracted at the stage for cotton plant fibrous cells to accumulate cellulose, and cDNA's complementary thereto were synthesized to construct a cDNA library. Clones of a number of 750 were arbitrarily selected from the library, and they were randomly subjected from to sequencing. Those having homology to an amino acid sequence deduced from a gene of cellulose 4-β-glucosyltransferase (bcsA) of cellulose synthase operon of acetic acid bacterium were selected from obtained nucleotide sequences of the respective clones. Thus, DNA coding for cellulose synthase was obtained.

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Description

Technical Field

The present invention relates to a DNA coding for cellulose synthase originating from cotton plant (Gossypium hirsutum), a recombinant DNA containing the same, a transformed cell transformed with the DNA, and a method for controlling cellular cellulose synthesis.

Background Art

Cellulose is used for paper, woody structural materials, fiber, cloths, food, cosmetics, and pharmaceuticals, as well [redacted] in biosynthesis of cellulose. The cellulose-related industry has been hitherto directed to such cellulose products that have been already produced. in which there has been no trial to develop a new material based on an aspect of biosynthesis. The mechanism of disease action, which is exerted by pathogenic microorganisms on plants, often results from the inhibition on cellulose biosynthesis as in Pyricularia oryzae (P. oryzae). Therefore, the addition of disease resistance to the cellulose biosynthesis mechanism is agriculturally applicable and valuable. Further, cellulose is the most abundant organic compound on the earth, and it is a sink in which the largest amount of CO_2 in the atmospheric air is fixed. Therefore, the genetic improvement of cellulose biosynthesis enzymes is also applicable to the industry which is directed to the control of CO_2 in the atmospheric air based on the use of cellulose as the sink.

In recent years, cDNA's originating from fiber cells of cotton plant have been randomly sequenced, and it has been reported that full length CelA1 and partial length of CelA2 probably represent cDNAs of cotton plant cellulose synthase, in view of the homology to bacterial cellulose synthase gene (bacterial BcsA) (Pear et al., Proceeding of National Academy of Science, USA (1996) 93 12637-12642). The binding ability to UDP-glucose has been demonstrated for CelA1. However, as for CelA2, the homology has been merely demonstrated for the C-terminal amino acid sequence.

Disclosure of the Invention

The present invention has been made in order to provide a new method for regulating cellulose production in prokaryotic cells or eukaryotic cells, an object of which is to provide a DNA coding for cellulose synthase, a recombinant DNA containing the same, a transformed cell transformed with the DNA, and a method for regulating cellular cellulose synthesis.

The present inventors firstly extracted mRNAs at the stage for cotton plant fiber cells to accumulate cellulose, and cDNAs complementary thereto were synthesized to construct a cDNA library. 750 of cDNA clones were arbitrarily selected from the library, and they were randomly subjected to sequencing. Six amino acid sequences were derived for one nucleotide sequence of each of the obtained clones to select those having homology to an amino acid sequence obtained by translation from a gene of cellulose 4-β-glucosyltransferase (bcsA) of cellulose synthase operon of acetobacterium. As a result, genes, which were classified into three types or groups, were found, and they were designated as PcsA1, PcsA2, and PcsA3 respectively (PcsA is an abbreviation of "Plant Cellulose Synthase A").

That is, the present invention lies in a DNA coding for any one of the following proteins (A) to (C):

(A) a protein having a cellulose synthase activity and comprising an amino acid sequence shown in SEQ ID NO: 2 or an amino acid sequence involving deletion, substitution, insertion, or addition of one or several amino acids relevant to SEQ ID NO: 2;

(B) a protein having a cellulose synthase activity and comprising an amino acid sequence shown in SEQ ID NO: 4 or an amino acid sequence involving deletion, substitution, insertion, or addition of one or several amino acids relevant to SEQ ID NO: 4: and

(C) a protein having a cellulose synthase activity and comprising an amino acid sequence shown in SEQ ID NO: 8 or an amino acid sequence involving deletion, substitution, insertion, or addition of one or several amino acids relevant to SEQ ID NO: 8, and comprising an amino acid sequence shown in SEQ ID NO: 11 or an amino acid sequence involving deletion, substitution, insertion, or addition of one or several amino acids relevant to SEQ ID NO: 11.

In another aspect, the present invention provides a recombinant vector comprising all or a part of the DNA as defined above, and a transformed cell transformed with the DNA as defined above.

In still another aspect, the present invention provides a method for regulating cellulose synthesis in a cell, comprising the steps of introducing the DNA as defined above into the cell, and expressing RNA having a nucleotide

sequence homologous to the DNA as defined above or a nucleotide sequence complementary to the DNA as define above.

SEQ ID NO: 1 corresponds to a sequence of PcsA1, and SEQ ID NO: 3 corresponds to a sequence of PcsA2. SEQ ID NO: 5 corresponds to a sequence of 3'-side region of PcsA3, SEQ ID NO: 7 corresponds to a sequence of 5'-side region of PcsA3, and SEQ ID NO: 9 corresponds to a sequence of internal region of PcsA3.

It has been demonstrated that PcsA1 and PcsA2 of the DNA's described above are DNA's coding for cotton plant cellulose synthase, according to the expression in eukaryotic cells (animal cells and/or yeast). It has been also demonstrated that an antibody thereagainst inhibits the cotton plant cellulose synthase activity in a cell-free system. Further, PcsA3, which is different from PcsA1 and PcsA2, has been found. Any one of these species was obtained as partial one, at the stage of clones obtained by the random sequencing, and no 5'-portion of the coding region was contained. Therefore, clones which have sequences of 5'-portions were isolated in accordance with the 5'-RACE method based on the use of PCR to determine the sequences. As a result of this operation, the sequences of the 5'-portions corresponding to the partial length clones were obtained for PcsA1 and PcsA2.

On the other hand, as for PcsA3, a sequence of a 5'-portion of another clone, which was considered to belong to the same PcsA3 group, was obtained. The both sequences had extremely high homology, and hence they were considered to have underwent multiple gene formation relatively recently originating from an identical gene through the process of duplication. Therefore, even when the both are combined with each other at corresponding portions to construct a fused gene followed by expression, it is assumed that the activity and function of a produced enzyme may not be affected thereby.

As for PcsA1 and PcsA2, in order to obtain a full length clone, primers were designed on the basis of the sequence of the 5'-portion and the sequence of the 3'-portion of the partial length clone to perform PCR. Thus, a clone containing ORF was obtained.

Those applicable as the template to be used for the RACE method may be any of cDNA synthesized from mRNA and a phage library. When the phage library is used, it is possible to use a sequence in the vector as a 5'-side primer.

As a result of random sequencing, seven clones concerning PcsA2 were most abundantly present, of 15 clones seemed to code the cellulose synthase. Expression was confirmed in eukaryotic cells (animal cells and/or yeast) transformed with the cellulose synthase gene. As a result, the cellulose synthase activity was observed.

The present invention will be explained in detail below.

<1> Preparation of cotton plant cDNA library

Cotton plant fiber cells at the stage of cellulose accumulation are preferably used as a material for extracting mRNA to construct a cotton plant cDNA library. The method for extracting mRNA is not specifically limited, for which it is possible to adopt an ordinary method for extracting mRNA from plant.

cDNA can be synthesized, for example, by using a poly T sequence which is complementary to poly A nucleotide existing at the terminal of mRNA as a primer to synthesize complementary DNA by the aid of reverse transcriptase, and forming a double strand by the aid of DNA polymerase.

The method therefor is described, for example, in Molecular Cloning (Maniatis et al., Cold Spring Harbour Laboratory). However, a variety of cDNA synthesis kits are commercially available from various companies, which may be used.

Generally, the library is constructed by using a phage vector. A variety of commercially available vectors are usable. However, it is preferable to use a vector, for example, λZAP vector in which it is unnecessary to perform recloning from the vector, and it is possible to immediately prepare a plasmid for sequencing.

<2> Determination of nucleotide sequence of cDNA

Clones are randomly selected from the obtained cDNA library to determine nucleotide sequences of inserts in the clones. The nucleotide sequence can be determined in accordance with the Maxam-Gilbert method or the dideoxy method. Among them, the dideoxy method is more convenient and preferred.

The nucleotide sequence can be determined in accordance with the dideoxy method by using a commercially available sequencing kit. Further, the use of an automatic sequencer makes it possible to determine sequences of a large number of clones for a short period of time.

It is unnecessary to determine the sequence for an entire length of the insert. It is enough to determine a length of nucleotide sequence which is considered to be sufficient to perform homology search. For example, in Examples described later on, the homology search as described below was performed when a sequence having not less than 60 nucleotides was successfully determined.

<3> Homology search with gene data base

The determined nucleotide sequence of each of cDNA clones is used to perform the homology search with respect to known amino acid sequences of the cellulose synthase or nucleotide sequences of genes coding therefor registered in the gene data base. The cellulose synthase is exemplified by an enzyme encoded by a gene of cellulose 4-β-glucosyltransferase (BcsA) of cellulose synthase operon of acetobacterium (Wong, H. C. et al., Proc. Natl. Acad. Sci. U.S.A., 87, 8130-8134 (1990). ACCESSION No. M37202).

Those usable as the data base include, for example, GenBank, EMBL, and DDBJ published, for example, from Los Alamos National Institute in the United States, Institute of European Molecular Biology, and National Institute of Genetics (Japan). Those commercially available and useable as the program for homology search include, for example, commercially available DNA analysis softwares, such as DNASIS (Hitachi Software Engineering Co.,Ltd.) and GENE-[redacted] nected on Internet with NCBI (National Center for Biotechnology Information) to utilize (http://www.ncbi.nlm.nih.gov/BLAST/) BLAST (Basic Local Alignment Search Tool) so that high speed homology search is performed.

The homology search is performed, for example, in accordance with the following algorithm. When the homology search is performed for a nucleotide sequence, homology comparison is advanced while shifting the nucleotide sequence to be investigated by every one nucleotide with respect to individual gene sequences included in the data base. When six or more continuous nucleotides are coincident, the homology score is counted and calculated in accordance with a homology score table (see, for example, M. Dayhoff, Atlas of Protein Sequence and Structure, vol. 5 (1978)). The system is set so that those having a score not less than a certain value are picked up as candidates which have homology. Further, the gap may be introduced into the sequence to be investigated or into the gene sequence included in the data base to make optimization so that the score is maximized.

When the homology search is performed for an amino acid sequence, a nucleotide sequence to be investigated is converted into amino acids concerning all six frames including those of a complementary chain. The investigation may be performed in the same manner as performed for the nucleotide. Specifically, it is possible to use blastx of BLAST described above. As for detailed techniques and conditions for the search, reference may be made to DDBJ News Letter, No. 15 (February 1995).

<4> Isolation of cDNA clone of cotton plant cellulose synthase

The clone obtained as described above is not necessarily contain the entire nucleotide sequence of the gene. In such a case, the clone is used as a probe to perform screening by means of plaque hybridization. Thus, it is possible to obtain a clone containing a full length gene from the library. A specified method may be carried out with reference to Molecular Cloning, second edition (Maniatis et al., Cold Spring Harbour Laboratory) 12.30 to 12.40.

When obtained cDNA is deficient in 5'-portion, the 5'-portion can be obtained as well by synthesizing primers so that the cDNA sequence may be elongated toward the 5'-terminal, and performing RT-PCR by using mRNA as a template.

As demonstrated in Examples described later on, the DNA of the present invention has been obtained as those having homology to the known bacterial cellulose synthase gene. The DNA further codes for an amino acid sequence GlnXXXXXXArgTrp (SEQ ID NO: 12) which is considered to form a UDP-glucose binding domain, having high homology in the vicinity thereof.

The nucleotide sequences of DNA of the present invention obtained as described above and the amino acid sequences deduced from the nucleotide sequences are shown in SEQ ID NOs: 1 to 10 in Sequence Listing. SEQ ID NOs: 1 and 3 show nucleotide sequences of PcsA1 and PcsA2 respectively. SEQ ID NOs: 2 and 4 show amino acid sequences deduced from the nucleotide sequences of PcsA1 and PcsA2 respectively.

SEQ ID NOs: 5 and 6 show a nucleotide sequence of a clone (PcsA3-682) containing 3'-side region of PcsA3 and an amino acid sequence deduced from the nucleotide sequence respectively. SEQ ID NOs: 7 and 8 show a nucleotide sequence of a 5'-portion (PcsA3-5') of another clone containing 5'-side region of PcsA3 and an amino acid sequence deduced from the nucleotide sequence respectively. SEQ ID NOs: 9 and 10 show a nucleotide sequence of 3'-portion (PcsA3-3') of the clone and an amino acid sequence deduced from the nucleotide sequence respectively (see Fig. 1). That is, SEQ ID NO: 5 corresponds to the 3'-side region of PcsA3, SEQ ID NO: 7 corresponds to the 5'-side region of PcsA3, and SEQ ID NO: 9 corresponds to internal region of PcsA3. The overlapping portion of PcsA3-682 is different from that of PcsA3-3' in 9 nucleotides in the nucleotide sequence and 1 amino acid in the amino acid sequence. Figs. 3 and 4 show the comparison between the nucleotide sequences of PcsA3-682 and PcsA3-3'. SEQ ID NO: 11 shows a combination of the amino acid sequences encoded by PcsA3-682 and PcsA3-3'.

The sequence of GlnXXXXXXArgTrp (SEQ ID NO: 12) corresponds to amino acid numbers 710 to 714 in SEQ ID NO: 2 for PcsA1, amino acid numbers 778 to 782 in SEQ ID NO: 4 for PcsA2, and amino acid numbers 356 to 360 in

530 535 540
His Asp Arg Tyr Ala Asn Arg Asn Val Val Phe Phe Asp Ile Asn Met
545 550 555 560
Leu Gly Leu Asp Gly Leu Gln Gly Pro Val Tyr Val Gly Thr Gly Cys
565 570 575
Val Phe Asn Arg Gln Ala Leu Tyr Gly Tyr Asp Pro Pro Val Ser Glu
580 585 590
Lys Arg Pro Lys Met Thr Cys Asp Cys Trp Pro Ser Trp Cys Cys Cys
595 600 605
Cys Cys Gly Gly Ser Arg Lys Lys Ser Lys Lys Lys Gly Glu Lys Lys
610 615 620
Gly Leu Leu Gly Gly Leu Leu Tyr Gly Lys Lys Lys Lys Met Met Gly
625 630 635 640
Lys Asn Tyr Val Lys Lys Gly Ser Ala Pro Val Phe Asp Leu Glu Glu
645 650 655
Ile Glu Glu Gly Leu Glu Gly Tyr Glu Glu Leu Glu Lys Ser Thr Leu
660 665 670
Met Ser Gln Lys Asn Phe Glu Lys Arg Phe Gly Gln Ser Pro Val Phe
675 680 685
Ile Ala Ser Thr Leu Met Glu Asn Gly Gly Leu Pro Glu Gly Thr Asn
690 695 700
Ser Thr Ser Leu Ile Lys Glu Ala Ile His Val Ile Ser Cys Gly Tyr
705 710 715 720
Glu Glu Lys Thr Glu Trp Gly Lys Glu Ile Gly Trp Ile Tyr Gly Ser
725 730 735
Val Thr Glu Asp Ile Leu Thr Gly Phe Lys Met His Cys Arg Gly Trp
740 745 750
Lys Ser Val Tyr Cys Val Pro Lys Arg Pro Ala Phe Lys Gly Ser Ala
755 760 765
Pro Ile Asn Leu Ser Asp Arg Leu His Gln Val Leu Arg Trp Ala Leu
770 775 780
Gly Ser Val Glu Ile Phe Leu Ser Arg His Cys Pro Leu Trp Tyr Gly
785 790 795 800
Tyr Gly Gly Lys Leu Lys Trp Leu Glu Arg Leu Ala Tyr Ile Asn Thr
805 810 815
Ile Val Tyr Pro Phe Thr Ser Ile Pro Leu Leu Ala Tyr Cys Thr Ile
820 825 830
Pro Ala Val Cys Leu Leu Thr Gly Lys Phe Ile Ile Pro Thr Leu Ser
835 840 845
Asn Leu Thr Ser Val Trp Phe Leu Ala Leu Phe Leu Ser Ile Ile Ala
850 855 860

Thr Gly Val Leu Glu Leu Arg Trp Ser Gly Val Ser Ile Gln Asp Trp
865 870 875 880
Trp Arg Asn Glu Gln Phe Trp Val Ile Gly Gly Val Ser Ala His Leu
885 890 895
Phe Ala Val Phe Gln Gly Leu Leu Lys Val Leu Ala Gly Val Asp Thr
900 905 910
Asn Phe Thr Val Thr Ala Lys Ala Ala Asp Asp Thr Glu Phe Gly Glu
915 920 925

[redacted]

Ile Ile Leu Asn Met Val Gly Val Val Ala Gly Val Ser Asp Ala Ile
945 950 955 960
Asn Asn Gly Tyr Gly Ser Trp Gly Pro Leu Phe Gly Lys Leu Phe Phe
965 970 975
Ala Phe Trp Val Ile Leu His Leu Tyr Pro Phe Leu Lys Gly Leu Met
980 985 990
Gly Arg Gln Asn Arg Thr Pro Thr Ile Val Val Leu Trp Ser Ile Leu
995 1000 1005
Leu Ala Ser Ile Phe Ser Leu Val Trp Val Arg Ile Asp Pro Phe Leu
1010 1015 1020
Pro Lys Gln Thr Gly Pro Val Leu Lys Gln Cys Gly Val Glu Cys
1025 1030 1035

(2) INFORMATION FOR SEQ ID NO: 5:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2033 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA to mRNA

(vi) ORIGINAL SOURCE:

(A) ORGANISM: Gossypium hirsutum L.

(C) INDIVIDUAL ISOLATE: Coker312

(ix) FEATURE:

(A) NAME/KEY: CDS

(B) LOCATION:1..1857

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 5:

CCG ACA TTC GTG AAG GAG CGT CGA GCT ATG AAG AGA GAA TAT GAA GAA 48
Pro Thr Phe Val Lys Glu Arg Arg Ala Met Lys Arg Glu Tyr Glu Glu
1 5 10 15
TTC AAG GTT AGG ATA AAT GCA CTT GTA GCC AAA GCC CAA AAG GTT CCT 96

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Phe Lys Val Arg Ile Asn Ala Leu Val Ala Lys Ala Gln Lys Val Pro
                20                  25                  30
CCA GAA GGG TGG ATC ATG CAA GAT GGG ACA CCA TGG CCA GGA AAC AAT   144
Pro Glu Gly Trp Ile Met Gln Asp Gly Thr Pro Trp Pro Gly Asn Asn
            35                  40                  45
ACT AAA GAT CAC CCT GGT ATG ATT CAA GTA TTT CTC GGT CAA AGT GGA   192
Thr Lys Asp His Pro Gly Met Ile Gln Val Phe Leu Gly Gln Ser Gly
        50                  55                  60
GGC CAT GAT ACC GAA GGA AAT GAG CTT CCT CGT CTC GTC TAT GTA TCT   240
Gly His Asp Thr Glu Gly Asn Glu Leu Pro Arg Leu Val Tyr Val Ser
    65                  70                  75                  80
CGA GAG AAA AGG CCT GGT TTC TTG CAT CAC AAG AAA GCT GGT GCC ATG   288
Arg Glu Lys Arg Pro Gly Phe Leu His His Lys Lys Ala Gly Ala Met
                85                  90                  95
AAC GCC CTT GTT CGG GTC TCG GGG GTG CTC ACA AAT GCT CCT TTT ATG   336
Asn Ala Leu Val Arg Val Ser Gly Val Leu Thr Asn Ala Pro Phe Met
            100                 105                 110
TTG AAC TTG GAT TGT GAC CAT TAT TTA AAT AAC AGC AAG GCT GTA AGA   384
Leu Asn Leu Asp Cys Asp His Tyr Leu Asn Asn Ser Lys Ala Val Arg
        115                 120                 125
GAG GCT ATG TGT TTC TTG ATG GAC CCT CAA ATT GGA AGA AAG GTT TGC   432
Glu Ala Met Cys Phe Leu Met Asp Pro Gln Ile Gly Arg Lys Val Cys
    130                 135                 140
TAT GTC CAA TTC CCT CAA CGT TTC GAT GGT ATT GAT AGA CAT GAT CGA   480
Tyr Val Gln Phe Pro Gln Arg Phe Asp Gly Ile Asp Arg His Asp Arg
145                 150                 155                 160
TAT GCC AAT CGG AAC ACA GTT TTC TTT GAT ATT AAC ATG AAA GGT CTA   528
Tyr Ala Asn Arg Asn Thr Val Phe Phe Asp Ile Asn Met Lys Gly Leu
                165                 170                 175
GAT GGT ATA CAA GGC CCT GTA TAT GTC GGC ACG GGG TGT GTT TTC AGA   576
Asp Gly Ile Gln Gly Pro Val Tyr Val Gly Thr Gly Cys Val Phe Arg
            180                 185                 190
AGG CAA GCT CTT TAT GGT TAT GAA CCT CCA AAG GGA CCT AAG CGC CCG   624
Arg Gln Ala Leu Tyr Gly Tyr Glu Pro Pro Lys Gly Pro Lys Arg Pro
        195                 200                 205
AAA ATG GTA ACC TGT GGT TGC TGC CCT TGT TTT GGA CGC CGC AGA AAG   672
Lys Met Val Thr Cys Gly Cys Cys Pro Cys Phe Gly Arg Arg Arg Lys
    210                 215                 220
GAC AAA AAG CAC TCT AAG GAT GGT GGA AAT GCA AAT GGT CTA AGC CTA   720
Asp Lys Lys His Ser Lys Asp Gly Gly Asn Ala Asn Gly Leu Ser Leu
225                 230                 235                 240
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GAA GCA GCC AAA GAT GAC AAG GAG TTA TTG ATG TCC CAC ATG AAC TTT        768
Glu Ala Ala Lys Asp Asp Lys Glu Leu Leu Met Ser His Met Asn Phe
                245                 250                 255
GAA AAG AAA TTT GGA CAA TCA GCC ATT TTT GTA ACT TCA ACA CTG ATG        816
Glu Lys Lys Phe Gly Gln Ser Ala Ile Phe Val Thr Ser Thr Leu Met
            260                 265                 270
GAA CAA GGT GGT GTC CCT CCT TCT TCA AGC CCC GCA GCT TTG CTC AAA        864
Glu Gln Gly Gly Val Pro Pro Ser Ser Ser Pro Ala Ala Leu Leu Lys
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[redacted]

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Glu Ala Ile His Val Ile Ser Cys Gly Tyr Glu Asp Lys Thr Glu Trp
    290                 295                 300
GGA AGC GAG CTT GGC TGG ATT TAC GGC TCG ATT ACA GAA GAT ATC TTA        960
Gly Ser Glu Leu Gly Trp Ile Tyr Gly Ser Ile Thr Glu Asp Ile Leu
305                 310                 315                 320
ACA GGA TTC AAG ATG CAT TGC CGT GGA TGG AGA TCA ATA TAC TGC ATG       1008
Thr Gly Phe Lys Met His Cys Arg Gly Trp Arg Ser Ile Tyr Cys Met
                325                 330                 335
CCA AAG TTG CCT GCA TTC AAG GGT TCA GCT CCC ATC AAT CTA TCG GAT       1056
Pro Lys Leu Pro Ala Phe Lys Gly Ser Ala Pro Ile Asn Leu Ser Asp
            340                 345                 350
CGT CTA AAC CAA GTC CTT CGA TGG GCA CTC GGT TCT GTT GAA ATT TTC       1104
Arg Leu Asn Gln Val Leu Arg Trp Ala Leu Gly Ser Val Glu Ile Phe
        355                 360                 365
TTT AGT CAT CAT TGC CCA GCA TGG TAT GGT TTC AAG GGA GGA AAG CTA       1152
Phe Ser His His Cys Pro Ala Trp Tyr Gly Phe Lys Gly Gly Lys Leu
    370                 375                 380
AAA TGG CTT GAA CGA TTC GCA TAT GTC AAC ACA ACC ATC TAC CCC TTC       1200
Lys Trp Leu Glu Arg Phe Ala Tyr Val Asn Thr Thr Ile Tyr Pro Phe
385                 390                 395                 400
ACA TCT TTA CCA CTT CTC GCC TAT TGT ACC CTA CCG GCA ATC TGT TTA       1248
Thr Ser Leu Pro Leu Leu Ala Tyr Cys Thr Leu Pro Ala Ile Cys Leu
                405                 410                 415
CTT ACC GAT AAA TTT ATC ATG CCA CCG ATA AGC ACC TTT GCA AGT CTA       1296
Leu Thr Asp Lys Phe Ile Met Pro Pro Ile Ser Thr Phe Ala Ser Leu
            420                 425                 430
TTC TTC ATT GCC TTG TTT CTT TCA ATC TTT GCA ACT GGT ATT CTC GAG       1344
Phe Phe Ile Ala Leu Phe Leu Ser Ile Phe Ala Thr Gly Ile Leu Glu
        435                 440                 445
CTA AGG TGG AGT GGA GTA AGC ATT GAA GAA TGG TGG AGG AAT GAG CAA       1392
Leu Arg Trp Ser Gly Val Ser Ile Glu Glu Trp Trp Arg Asn Glu Gln
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     450                 455                 460
TTT TGG GTC ATC GGT GGC ATT TCG GCA CAT TTG TTC GCT GTT ATC CAA      1440
Phe Trp Val Ile Gly Gly Ile Ser Ala His Leu Phe Ala Val Ile Gln
465                 470                 475                 480
GGC TTG TTG AAA GTT CTA GCT GGT ATT GAC ACT AAT TTC ACT GTC ACA      1488
Gly Leu Leu Lys Val Leu Ala Gly Ile Asp Thr Asn Phe Thr Val Thr
                485                 490                 495
TCC AAG GCA ACT GAT GAC GAG GAG TTC GGG GAA TTG TAT ACT TTC AAA      1536
Ser Lys Ala Thr Asp Asp Glu Glu Phe Gly Glu Leu Tyr Thr Phe Lys
            500                 505                 510
TGG ACA ACC CTT CTA ATT CCT CCT ACT ACC GTC TTA ATC ATC AAT TTA      1584
Trp Thr Thr Leu Leu Ile Pro Pro Thr Thr Val Leu Ile Ile Asn Leu
        515                 520                 525
GTC GGT GTC GTT GCA GGC ATC TCG GAT GCC ATA AAC AAT GGA TAC CAA      1632
Val Gly Val Val Ala Gly Ile Ser Asp Ala Ile Asn Asn Gly Tyr Gln
    530                 535                 540
TCA TGG GGA CCT CTT TTT GGG AAG CTC TTC TTC TCT TTC TGG GTG ATT      1680
Ser Trp Gly Pro Leu Phe Gly Lys Leu Phe Phe Ser Phe Trp Val Ile
545                 550                 555                 560
GTC CAT CTC TAT CCA TTC CTC AAA GGT TTA ATG GGG AGA CAA AAC CGG      1728
Val His Leu Tyr Pro Phe Leu Lys Gly Leu Met Gly Arg Gln Asn Arg
                565                 570                 575
ACA CCA ACC ATT GTT GTT ATA TGG TCA GTG CTA TTG GCT TCA ATC TTC      1776
Thr Pro Thr Ile Val Val Ile Trp Ser Val Leu Leu Ala Ser Ile Phe
            580                 585                 590
TCC TTG CTT TGG GTC CGA ATT GAT CCA TTT GTG ATG AAA ACC AAA GGA      1824
Ser Leu Leu Trp Val Arg Ile Asp Pro Phe Val Met Lys Thr Lys Gly
        595                 600                 605
CCA GAC ACT ACA ATG TGT GGC ATT AAC TGT TGAAAAAAAA TCATCTTGCG        1874
Pro Asp Thr Thr Met Cys Gly Ile Asn Cys
    610                 615
TGGTTCTTTT AGATTATGGT ATGTGATGTA TGAACAAACA AGAATGGAGA TGCACAAGAC    1934
AGAATAAAAT TAGAGTGAAA GTTTTGTGTA GTTATATATT CATTCTACCA ACTATAAGTT    1994
TTGTCATTCA ATTGAAAATA GCTCAACTTT GTGATCAAA                           2033
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(2) INFORMATION FOR SEQ ID NO: 6:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 618 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(v) FRAGMENT TYPE: C-terminal fragment

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 6:

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Pro Thr Phe Val Lys Glu Arg Arg Ala Met Lys Arg Glu Tyr Glu Glu
 1               5                  10                  15
Phe Lys Val Arg Ile Asn Ala Leu Val Ala Lys Ala Gln Lys Val Pro
            20                  25                  30
Pro Glu Gly Trp Ile Met Gln Asp Gly Thr Pro Trp Pro Gly Asn Asn
        35                  40                  45
[redacted]
    50                  55                  60
Gly His Asp Thr Glu Gly Asn Glu Leu Pro Arg Leu Val Tyr Val Ser
65                  70                  75                  80
Arg Glu Lys Arg Pro Gly Phe Leu His His Lys Lys Ala Gly Ala Met
                85                  90                  95
Asn Ala Leu Val Arg Val Ser Gly Val Leu Thr Asn Ala Pro Phe Met
            100                 105                 110
Leu Asn Leu Asp Cys Asp His Tyr Leu Asn Asn Ser Lys Ala Val Arg
        115                 120                 125
Glu Ala Met Cys Phe Leu Met Asp Pro Gln Ile Gly Arg Lys Val Cys
    130                 135                 140
Tyr Val Gln Phe Pro Gln Arg Phe Asp Gly Ile Asp Arg His Asp Arg
145                 150                 155                 160
Tyr Ala Asn Arg Asn Thr Val Phe Phe Asp Ile Asn Met Lys Gly Leu
                165                 170                 175
Asp Gly Ile Gln Gly Pro Val Tyr Val Gly Thr Gly Cys Val Phe Arg
            180                 185                 190
Arg Gln Ala Leu Tyr Gly Tyr Glu Pro Pro Lys Gly Pro Lys Arg Pro
        195                 200                 205
Lys Met Val Thr Cys Gly Cys Cys Pro Cys Phe Gly Arg Arg Arg Lys
    210                 215                 220
Asp Lys Lys His Ser Lys Asp Gly Gly Asn Ala Asn Gly Leu Ser Leu
225                 230                 235                 240
Glu Ala Ala Lys Asp Asp Lys Glu Leu Leu Met Ser His Met Asn Phe
                245                 250                 255
Glu Lys Lys Phe Gly Gln Ser Ala Ile Phe Val Thr Ser Thr Leu Met
            260                 265                 270
Glu Gln Gly Gly Val Pro Pro Ser Ser Ser Pro Ala Ala Leu Leu Lys
        275                 280                 285
Glu Ala Ile His Val Ile Ser Cys Gly Tyr Glu Asp Lys Thr Glu Trp
    290                 295                 300
Gly Ser Glu Leu Gly Trp Il  Tyr Gly Ser Ile Thr Glu Asp Ile Leu
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305 310 315 320
Thr Gly Phe Lys Met His Cys Arg Gly Trp Arg Ser Ile Tyr Cys Met
325 330 335
Pro Lys Leu Pro Ala Phe Lys Gly Ser Ala Pro Ile Asn Leu Ser Asp
340 345 350
Arg Leu Asn Gln Val Leu Arg Trp Ala Leu Gly Ser Val Glu Ile Phe
355 360 365
Phe Ser His His Cys Pro Ala Trp Tyr Gly Phe Lys Gly Gly Lys Leu
370 375 380
Lys Trp Leu Glu Arg Phe Ala Tyr Val Asn Thr Thr Ile Tyr Pro Phe
385 390 395 400
Thr Ser Leu Pro Leu Leu Ala Tyr Cys Thr Leu Pro Ala Ile Cys Leu
405 410 415
Leu Thr Asp Lys Phe Ile Met Pro Pro Ile Ser Thr Phe Ala Ser Leu
420 425 430
Phe Phe Ile Ala Leu Phe Leu Ser Ile Phe Ala Thr Gly Ile Leu Glu
435 440 445
Leu Arg Trp Ser Gly Val Ser Ile Glu Glu Trp Trp Arg Asn Glu Gln
450 455 460
Phe Trp Val Ile Gly Gly Ile Ser Ala His Leu Phe Ala Val Ile Gln
465 470 475 480
Gly Leu Leu Lys Val Leu Ala Gly Ile Asp Thr Asn Phe Thr Val Thr
485 490 495
Ser Lys Ala Thr Asp Asp Glu Glu Phe Gly Glu Leu Tyr Thr Phe Lys
500 505 510
Trp Thr Thr Leu Leu Ile Pro Pro Thr Thr Val Leu Ile Ile Asn Leu
515 520 525
Val Gly Val Val Ala Gly Ile Ser Asp Ala Ile Asn Asn Gly Tyr Gln
530 535 540
Ser Trp Gly Pro Leu Phe Gly Lys Leu Phe Phe Ser Phe Trp Val Ile
545 550 555 560
Val His Leu Tyr Pro Phe Leu Lys Gly Leu Met Gly Arg Gln Asn Arg
565 570 575
Thr Pro Thr Ile Val Val Ile Trp Ser Val Leu Leu Ala Ser Ile Phe
580 585 590
Ser Leu Leu Trp Val Arg Ile Asp Pro Phe Val Met Lys Thr Lys Gly
595 600 605
Pro Asp Thr Thr Met Cys Gly Ile Asn Cys
610 615

(2) INFORMATION FOR SEQ ID NO: 7:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1086 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear
(ii) MOLECULE TYPE: cDNA to mRNA
(vi) ORIGINAL SOURCE:
(A) ORGANISM: Gossypium hirsutum L.

[redacted]

(ix) FEATURE:
(A) NAME/KEY: CDS
(B) LOCATION:24..1086
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 7:

```
GGCACGAGCT TTCATATCCT CCA ATG GAA GCC AGC GCC GGA CTC GTT GCG          50
                          Met Glu Ala Ser Ala Gly Leu Val Ala
                            1               5
GGC TCT CAC AAC CGC AAT GAA CTT GTT GTC ATT CAT GGC CAT GAA GAG          98
Gly Ser His Asn Arg Asn Glu Leu Val Val Ile His Gly His Glu Glu
 10                  15                  20                  25
CCT AAA CCT CTG AAG AAC TTG GAT GGT CAA GTT TGT GAG ATT TGT GGT         146
Pro Lys Pro Leu Lys Asn Leu Asp Gly Gln Val Cys Glu Ile Cys Gly
                30                  35                  40
GAT GAA ATT GGG TTG ACG GTC GAT GGA GAT CTT TTC GTG GCC TGC AAC         194
Asp Glu Ile Gly Leu Thr Val Asp Gly Asp Leu Phe Val Ala Cys Asn
            45                  50                  55
GAG TGT GGT TTT CCA GTT TGT AGG CCT TGT TAT GAG TAT GAA AGG AGA         242
Glu Cys Gly Phe Pro Val Cys Arg Pro Cys Tyr Glu Tyr Glu Arg Arg
        60                  65                  70
GAA GGG AGT CAA CAA TGT CCT CAA TGC AAA ACT AGA TAC AAG CGT CTC         290
Glu Gly Ser Gln Gln Cys Pro Gln Cys Lys Thr Arg Tyr Lys Arg Leu
    75                  80                  85
AAG GGG AGT CCG AGG GTG GAG GGA GAT GAA GAT GAA GAG GAT GTG GAT         338
Lys Gly Ser Pro Arg Val Glu Gly Asp Glu Asp Glu Glu Asp Val Asp
 90                  95                  100                 105
GAT ATC GAA CAT GAA TTC AAC ATT GAT GAT GAA CAA AAC AAG TAT AGA         386
Asp Ile Glu His Glu Phe Asn Ile Asp Asp Glu Gln Asn Lys Tyr Arg
                110                 115                 120
AAT ATC GCT GAA TCG ATG CTT CAT GGA AAG ATG AGC TAC GGG AGA GGC         434
Asn Ile Ala Glu Ser Met Leu His Gly Lys Met Ser Tyr Gly Arg Gly
            125                 130                 135
CCT GAA GAC GAT GAA GGT TTG CAA ATC CCA CCC GGT TTA GCT GGT GTT         482
```

```
Pro Glu Asp Asp Glu Gly Leu Gln Ile Pro Pro Gly Leu Ala Gly Val
        140                 145                 150
CGA TCT CGG CCG GTG AGC GGG GAG TTC CCA ATA GGA AGC TCT CTT GCT      530
Arg Ser Arg Pro Val Ser Gly Glu Phe Pro Ile Gly Ser Ser Leu Ala
    155                 160                 165
TAT GGG GAA CAC ATG TCA AAT AAA CGA GTT CAT CCA TAT CCT ATG TCT      578
Tyr Gly Glu His Met Ser Asn Lys Arg Val His Pro Tyr Pro Met Ser
170                 175                 180                 185
GAA CCT GGA AGT GCA AGA TGG GAT GAA AAG AAA GAG GGA GGA TGG AGA      626
Glu Pro Gly Ser Ala Arg Trp Asp Glu Lys Lys Glu Gly Gly Trp Arg
                190                 195                 200
GAA AGG ATG GAT GAT TGG AAA ATG CAG CAA GGG AAT TTG GGT CCT GAA      674
Glu Arg Met Asp Asp Trp Lys Met Gln Gln Gly Asn Leu Gly Pro Glu
            205                 210                 215
CCT GAT GAT GCC TAT GAT GCT GAC ATG GCT ATG CTT GAT GAA GCT AGG      722
Pro Asp Asp Ala Tyr Asp Ala Asp Met Ala Met Leu Asp Glu Ala Arg
        220                 225                 230
CAG CCA TTG TCA AGG AAA GTG CCA ATT GCA TCG AGC AAA ATC AAT CCT      770
Gln Pro Leu Ser Arg Lys Val Pro Ile Ala Ser Ser Lys Ile Asn Pro
    235                 240                 245
TAT CGT ATG GTG ATT GTG GCT CGT CTA GTT ATC CTT GCT TTC TTT CTT      818
Tyr Arg Met Val Ile Val Ala Arg Leu Val Ile Leu Ala Phe Phe Leu
250                 255                 260                 265
CGC TAT CGG ATT TTG AAC CCG GTA CAT GAT GCA ATT GGG CTT TGG CTA      866
Arg Tyr Arg Ile Leu Asn Pro Val His Asp Ala Ile Gly Leu Trp Leu
                270                 275                 280
ACT TCT GTG ATC TGT GAA ATC TGG TTT GCC TTT TCA TGG ATC CTT GAT      914
Thr Ser Val Ile Cys Glu Ile Trp Phe Ala Phe Ser Trp Ile Leu Asp
            285                 290                 295
CAG TTC CCT AAA TGG TTC CCT ATT GAC CGC GAG ACG TAT CTC GAT CGC      962
Gln Phe Pro Lys Trp Phe Pro Ile Asp Arg Glu Thr Tyr Leu Asp Arg
        300                 305                 310
CTT TCC CTC AGG TAT GAG AGG GAA GGT GAG CCC AAC ATG CTT GCT TCT      1010
Leu Ser Leu Arg Tyr Glu Arg Glu Gly Glu Pro Asn Met Leu Ala Ser
    315                 320                 325
GTT GAT ATT TTT GTC AGT ACA GTG GAT CCA TTG AAG GGA CCT CCT CTA      1058
Val Asp Ile Phe Val Ser Thr Val Asp Pro Leu Lys Gly Pro Pro Leu
330                 335                 340                 345
GTA ACA GCG AAT ACA GTT CTA TCG ATC T                               1086
Val Thr Ala Asn Thr Val Leu Ser Ile
                350
```

(2) INFORMATION FOR SEQ ID NO: 8:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 354 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(v) FRAGMENT TYPE: N-terminal fragment

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 8:

[redacted]

```
 1               5                  10                  15
Leu Val Val Ile His Gly His Glu Glu Pro Lys Pro Leu Lys Asn Leu
            20                  25                  30
Asp Gly Gln Val Cys Glu Ile Cys Gly Asp Glu Ile Gly Leu Thr Val
        35                  40                  45
Asp Gly Asp Leu Phe Val Ala Cys Asn Glu Cys Gly Phe Pro Val Cys
    50                  55                  60
Arg Pro Cys Tyr Glu Tyr Glu Arg Arg Glu Gly Ser Gln Gln Cys Pro
65                  70                  75                  80
Gln Cys Lys Thr Arg Tyr Lys Arg Leu Lys Gly Ser Pro Arg Val Glu
                85                  90                  95
Gly Asp Glu Asp Glu Glu Asp Val Asp Asp Ile Glu His Glu Phe Asn
            100                 105                 110
Ile Asp Asp Glu Gln Asn Lys Tyr Arg Asn Ile Ala Glu Ser Met Leu
        115                 120                 125
His Gly Lys Met Ser Tyr Gly Arg Gly Pro Glu Asp Asp Glu Gly Leu
    130                 135                 140
Gln Ile Pro Pro Gly Leu Ala Gly Val Arg Ser Arg Pro Val Ser Gly
145                 150                 155                 160
Glu Phe Pro Ile Gly Ser Ser Leu Ala Tyr Gly Glu His Met Ser Asn
                165                 170                 175
Lys Arg Val His Pro Tyr Pro Met Ser Glu Pro Gly Ser Ala Arg Trp
            180                 185                 190
Asp Glu Lys Lys Glu Gly Gly Trp Arg Glu Arg Met Asp Asp Trp Lys
        195                 200                 205
Met Gln Gln Gly Asn Leu Gly Pro Glu Pro Asp Asp Ala Tyr Asp Ala
    210                 215                 220
Asp Met Ala Met Leu Asp Glu Ala Arg Gln Pro Leu Ser Arg Lys Val
225                 230                 235                 240
Pro Ile Ala Ser Ser Lys Ile Asn Pro Tyr Arg Met Val Ile Val Ala
                245                 250                 255
```

```
Arg Leu Val Ile Leu Ala Phe Phe Leu Arg Tyr Arg Ile Leu Asn Pro
            260                 265                 270
Val His Asp Ala Ile Gly Leu Trp Leu Thr Ser Val Ile Cys Glu Ile
        275                 280                 285
Trp Phe Ala Phe Ser Trp Ile Leu Asp Gln Phe Pro Lys Trp Phe Pro
    290                 295                 300
Ile Asp Arg Glu Thr Tyr Leu Asp Arg Leu Ser Leu Arg Tyr Glu Arg
305                 310                 315                 320
Glu Gly Glu Pro Asn Met Leu Ala Ser Val Asp Ile Phe Val Ser Thr
                325                 330                 335
Val Asp Pro Leu Lys Gly Pro Pro Leu Val Thr Ala Asn Thr Val Leu
            340                 345                 350
Ser Ile
```

(2) INFORMATION FOR SEQ ID NO: 9:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1000 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA to mRNA

(vi) ORIGINAL SOURCE:

(A) ORGANISM: Gossypium hirsutum L.

(C) INDIVIDUAL ISOLATE: Coker312

(ix) FEATURE:

(A) NAME/KEY: CDS

(B) LOCATION:1..1000

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 9:

```
GAC AAA GTC CGG CCG ACA TTC GTG AAG GAG CGT CGA GCT ATG AAG AGA      48
Asp Lys Val Arg Pro Thr Phe Val Lys Glu Arg Arg Ala Met Lys Arg
 1               5                  10                  15
GAA TAT GAA GAA TTC AAG GTT AGG ATA AAT GCA CTT GTA GCC AAA GCC      96
Glu Tyr Glu Glu Phe Lys Val Arg Ile Asn Ala Leu Val Ala Lys Ala
            20                  25                  30
CAA AAG GTT CCT CCA GAA GGG TGG ATC ATG CAA GAT GGG ACA CCA TGG     144
Gln Lys Val Pro Pro Glu Gly Trp Ile Met Gln Asp Gly Thr Pro Trp
        35                  40                  45
CCA GGA AAC AAT ACT AAA GAT CAC CCT GGT ATG ATT CAA GTA TTT CTC     192
Pro Gly Asn Asn Thr Lys Asp His Pro Gly Met Ile Gln Val Phe Leu
    50                  55                  60
```

```
GGT CAA AGT GGA GGC CAT GAT ACC GAA GGA AAT GAG CTT CCT CGT CTC        240
Gly Gln Ser Gly Gly His Asp Thr Glu Gly Asn Glu Leu Pro Arg Leu
 65                  70                  75                  80
GTC TAT GTA TCT CGA GAG AAA AGG CCA GGT TTC TTG CAT CAC AAG AAA        288
Val Tyr Val Ser Arg Glu Lys Arg Pro Gly Phe Leu His His Lys Lys
                 85                  90                  95
GCT GGT GCC ATG AAC GCC CTT GTT CGT GTC TCG GGG GTG CTT ACA AAT        336
Ala Gly Ala Met Asn Ala Leu Val Arg Val Ser Gly Val Leu Thr Asn
            100                 105                 110
[redacted]
GCT CCT TTT ATG TTG AAC TTG GAT TGT GAC CAC TAT TTA AAT AAC AGC        384
Ala Pro Phe Met Leu Asn Leu Asp Cys Asp His Tyr Leu Asn Asn Ser
        115                 120                 125
AAG GCT GTA AGA GAG GCT ATG TGT TTC TTG ATG GAC CCT CAA ATT GGA        432
Lys Ala Val Arg Glu Ala Met Cys Phe Leu Met Asp Pro Gln Ile Gly
    130                 135                 140
AGA AAG GTT TGC TAT GTC CAA TTC CCT CAA CGT TTC GAT GGT ATT GAT        480
Arg Lys Val Cys Tyr Val Gln Phe Pro Gln Arg Phe Asp Gly Ile Asp
145                 150                 155                 160
AGA CAT GAT CGA TAT GCC AAT CGG AAC ACA GTT TTC TTT GAT ATT AAC        528
Arg His Asp Arg Tyr Ala Asn Arg Asn Thr Val Phe Phe Asp Ile Asn
                165                 170                 175
ATG AAA GGT CTA GAT GGT ATA CAA GGC CCT GTA TAT GTC GGC ACG GGG        576
Met Lys Gly Leu Asp Gly Ile Gln Gly Pro Val Tyr Val Gly Thr Gly
            180                 185                 190
TGT GTT TTC AGA AGG CAA GCT CTT TAT GGT TAT GAA CCT CCA AAG GGA        624
Cys Val Phe Arg Arg Gln Ala Leu Tyr Gly Tyr Glu Pro Pro Lys Gly
        195                 200                 205
CCT AAG CGC CCG AAA ATG GTA ACC TGT GGT TGC TGC CCT TGC TTT GGA        672
Pro Lys Arg Pro Lys Met Val Thr Cys Gly Cys Cys Pro Cys Phe Gly
    210                 215                 220
CGC CGC AGA AAG GAC AAA AAG CAC TCT AAG GAT GGT GGA AAT GCA AAT        720
Arg Arg Arg Lys Asp Lys Lys His Ser Lys Asp Gly Gly Asn Ala Asn
225                 230                 235                 240
GGT CTA AGC CTA GAA GCA GCC GAA GAT GAC AAG GAG TTA TTG ATG TCC        768
Gly Leu Ser Leu Glu Ala Ala Glu Asp Asp Lys Glu Leu Leu Met Ser
                245                 250                 255
CAC ATG AAC TTT GAA AAG AAA TTT GGA CAA TCA GCC ATT TTT GTA ACT        816
His Met Asn Phe Glu Lys Lys Phe Gly Gln Ser Ala Ile Phe Val Thr
            260                 265                 270
TCA ACA CTG ATG GAA CAA GGT GGT GTC CCT CCT TCT TCA AGC CCT GCA        864
Ser Thr Leu Met Glu Gln Gly Gly Val Pro Pro Ser Ser Ser Pro Ala
```

```
          275                 280                 285
GCT TTG CTC AAA GAA GCC ATT CAT GTA ATT AGT TGT GGT TAT GAA GAC      912
Ala Leu Leu Lys Glu Ala Ile His Val Ile Ser Cys Gly Tyr Glu Asp
     290                 295                 300
AAA ACC GAA TGG GGA AGC GAG CTT GGC TGG ATT TAC GGC TCG ATT ACA      960
Lys Thr Glu Trp Gly Ser Glu Leu Gly Trp Ile Tyr Gly Ser Ile Thr
305                 310                 315                 320
GAA GAT ATC TTA ACA GGT TTC AAG ATG CAT TGC CGT GGA  T               1000
Glu Asp Ile Leu Thr Gly Phe Lys Met His Cys Arg Gly
                325                 330
```

(2) INFORMATION FOR SEQ ID NO: 10:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 333 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(v) FRAGMENT TYPE: internal fragment

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 10:

```
Asp Lys Val Arg Pro Thr Phe Val Lys Glu Arg Arg Ala Met Lys Arg
 1               5                  10                  15
Glu Tyr Glu Glu Phe Lys Val Arg Ile Asn Ala Leu Val Ala Lys Ala
            20                  25                  30
Gln Lys Val Pro Pro Glu Gly Trp Ile Met Gln Asp Gly Thr Pro Trp
        35                  40                  45
Pro Gly Asn Asn Thr Lys Asp His Pro Gly Met Ile Gln Val Phe Leu
    50                  55                  60
Gly Gln Ser Gly Gly His Asp Thr Glu Gly Asn Glu Leu Pro Arg Leu
65                  70                  75                  80
Val Tyr Val Ser Arg Glu Lys Arg Pro Gly Phe Leu His His Lys Lys
                85                  90                  95
Ala Gly Ala Met Asn Ala Leu Val Arg Val Ser Gly Val Leu Thr Asn
            100                 105                 110
Ala Pro Phe Met Leu Asn Leu Asp Cys Asp His Tyr Leu Asn Asn Ser
        115                 120                 125
Lys Ala Val Arg Glu Ala Met Cys Phe Leu Met Asp Pro Gln Ile Gly
    130                 135                 140
Arg Lys Val Cys Tyr Val Gln Phe Pro Gln Arg Phe Asp Gly Ile Asp
145                 150                 155                 160
Arg His Asp Arg Tyr Ala Asn Arg Asn Thr Val Phe Phe Asp Ile Asn
                165                 170                 175
```

Met Lys Gly Leu Asp Gly Ile Gln Gly Pro Val Tyr Val Gly Thr Gly
180 185 190
Cys Val Phe Arg Arg Gln Ala Leu Tyr Gly Tyr Glu Pro Pro Lys Gly
195 200 205
Pro Lys Arg Pro Lys Met Val Thr Cys Gly Cys Cys Pro Cys Phe Gly
210 215 220
Arg Arg Arg Lys Asp Lys Lys His Ser Lys Asp Gly Gly Asn Ala Asn
225 230 235 240

245 250 255

His Met Asn Phe Glu Lys Lys Phe Gly Gln Ser Ala Ile Phe Val Thr
260 265 270
Ser Thr Leu Met Glu Gln Gly Gly Val Pro Pro Ser Ser Ser Pro Ala
275 280 285
Ala Leu Leu Lys Glu Ala Ile His Val Ile Ser Cys Gly Tyr Glu Asp
290 295 300
Lys Thr Glu Trp Gly Ser Glu Leu Gly Trp Ile Tyr Gly Ser Ile Thr
305 310 315 320
Glu Asp Ile Leu Thr Gly Phe Lys Met His Cys Arg Gly
325 330

(2) INFORMATION FOR SEQ ID NO: 11:
(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 622 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
(ii) MOLECULE TYPE: peptide
(v) FRAGMENT TYPE: C-terminal fragment
(ix) FEATURE:
(A) NAME/KEY:
(B) LOCATION:
(D) OTHER INFORMATION: Xaa indicates Glu or Lys
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 11:
Asp Lys Val Arg Pro Thr Phe Val Lys Glu Arg Arg Ala Met Lys Arg
1 5 10 15
Glu Tyr Glu Glu Phe Lys Val Arg Ile Asn Ala Leu Val Ala Lys Ala
20 25 30
Gln Lys Val Pro Pro Glu Gly Trp Ile Met Gln Asp Gly Thr Pro Trp
35 40 45
Pro Gly Asn Asn Thr Lys Asp His Pro Gly Met Ile Gln Val Phe Leu
50 55 60

Gly Gln Ser Gly Gly His Asp Thr Glu Gly Asn Glu Leu Pro Arg Leu
65 70 75 80
Val Tyr Val Ser Arg Glu Lys Arg Pro Gly Phe Leu His His Lys Lys
85 90 95
Ala Gly Ala Met Asn Ala Leu Val Arg Val Ser Gly Val Leu Thr Asn
100 105 110
Ala Pro Phe Met Leu Asn Leu Asp Cys Asp His Tyr Leu Asn Asn Ser
115 120 125
Lys Ala Val Arg Glu Ala Met Cys Phe Leu Met Asp Pro Gln Ile Gly
130 135 140
Arg Lys Val Cys Tyr Val Gln Phe Pro Gln Arg Phe Asp Gly Ile Asp
145 150 155 160
Arg His Asp Arg Tyr Ala Asn Arg Asn Thr Val Phe Phe Asp Ile Asn
165 170 175
Met Lys Gly Leu Asp Gly Ile Gln Gly Pro Val Tyr Val Gly Thr Gly
180 185 190
Cys Val Phe Arg Arg Gln Ala Leu Tyr Gly Tyr Glu Pro Pro Lys Gly
195 200 205
Pro Lys Arg Pro Lys Met Val Thr Cys Gly Cys Cys Pro Cys Phe Gly
210 215 220
Arg Arg Arg Lys Asp Lys Lys His Ser Lys Asp Gly Gly Asn Ala Asn
225 230 235 240
Gly Leu Ser Leu Glu Ala Ala Xaa Asp Asp Lys Glu Leu Leu Met Ser
245 250 255
His Met Asn Phe Glu Lys Lys Phe Gly Gln Ser Ala Ile Phe Val Thr
260 265 270
Ser Thr Leu Met Glu Gln Gly Gly Val Pro Pro Ser Ser Ser Pro Ala
275 280 285
Ala Leu Leu Lys Glu Ala Ile His Val Ile Ser Cys Gly Tyr Glu Asp
290 295 300
Lys Thr Glu Trp Gly Ser Glu Leu Gly Trp Ile Tyr Gly Ser Ile Thr
305 310 315 320
Glu Asp Ile Leu Thr Gly Phe Lys Met His Cys Arg Gly Trp Arg Ser
325 330 335
Ile Tyr Cys Met Pro Lys Leu Pro Ala Phe Lys Gly Ser Ala Pro Ile
340 345 350
Asn Leu Ser Asp Arg Leu Asn Gln Val Leu Arg Trp Ala Leu Gly Ser
355 360 365
Val Glu Ile Phe Phe Ser His His Cys Pro Ala Trp Tyr Gly Phe Lys
370 375 380
Gly Gly Lys Leu Lys Trp Leu Glu Arg Phe Ala Tyr Val Asn Thr Thr

385 390 395 400
Ile Tyr Pro Phe Thr Ser Leu Pro Leu Leu Ala Tyr Cys Thr Leu Pro
405 410 415
Ala Ile Cys Leu Leu Thr Asp Lys Phe Ile Met Pro Pro Ile Ser Thr
420 425 430
Phe Ala Ser Leu Phe Phe Ile Ala Leu Phe Leu Ser Ile Phe Ala Thr
435 440 445
Gly Ile Leu Glu Leu Arg Trp Ser Gly Val Ser Ile Glu Glu Trp Trp

Arg Asn Glu Gln Phe Trp Val Ile Gly Gly Ile Ser Ala His Leu Phe
465 470 475 480
Ala Val Ile Gln Gly Leu Leu Lys Val Leu Ala Gly Ile Asp Thr Asn
485 490 495
Phe Thr Val Thr Ser Lys Ala Thr Asp Asp Glu Glu Phe Gly Glu Leu
500 505 510
Tyr Thr Phe Lys Trp Thr Thr Leu Leu Ile Pro Pro Thr Thr Val Leu
515 520 525
Ile Ile Asn Leu Val Gly Val Val Ala Gly Ile Ser Asp Ala Ile Asn
530 535 540
Asn Gly Tyr Gln Ser Trp Gly Pro Leu Phe Gly Lys Leu Phe Phe Ser
545 550 555 560
Phe Trp Val Ile Val His Leu Tyr Pro Phe Leu Lys Gly Leu Met Gly
565 570 575
Arg Gln Asn Arg Thr Pro Thr Ile Val Val Ile Trp Ser Val Leu Leu
580 585 590
Ala Ser Ile Phe Ser Leu Leu Trp Val Arg Ile Asp Pro Phe Val Met
595 600 605
Lys Thr Lys Gly Pro Asp Thr Thr Met Cys Gly Ile Asn Cys
610 615 620

(2) INFORMATION FOR SEQ ID NO: 12:
(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 9 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
(ii) MOLECULE TYPE: peptide
(v) FRAGMENT TYPE: internal fragment
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 12:
Gln Xaa Xaa Xaa Xaa Xaa Xaa Arg Trp
1 5

(2) INFORMATION FOR SEQ ID NO: 13:
(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 50 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear
(ii) MOLECULE TYPE: other nucleic acid
(A) DESCRIPTION: /desc = "Synthetic DNA"
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 13:
GAGAGAGAGA GAGAGAGAGA ACTAGTCTCG AGTTTTTTTT TTTTTTTTTT 50

(2) INFORMATION FOR SEQ ID NO: 14:
(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 13 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear
(ii) MOLECULE TYPE: other nucleic acid
(A) DESCRIPTION: /desc = "Synthetic DNA"
(ix) FEATURE:
(A) NAME/KEY:
(B) LOCATION:1..4
(D) OTHER INFORMATION: single strand
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 14:
AATTCGGCAC GAG 13

(2) INFORMATION FOR SEQ ID NO: 15:
(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 20 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear
(ii) MOLECULE TYPE: other nucleic acid
(A) DESCRIPTION: /desc = "Synthetic DNA"
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 15:
GACTGAAGAT AAGCCAAAAG 20

(2) INFORMATION FOR SEQ ID NO: 16:
(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 19 base pairs
(B) TYPE: nucleic acid

(C) STRANDEDNESS: single
(D) TOPOLOGY: linear
(ii) MOLECULE TYPE: other nucleic acid
(A) DESCRIPTION: /desc = "Synthetic DNA"
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 16:
GGAATGATGA ATTTGCCGG 19

(2) INFORMATION FOR SEQ ID NO: 17:

[redacted]

(A) LENGTH: 20 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear
(ii) MOLECULE TYPE: other nucleic acid
(A) DESCRIPTION: /desc = "Synthetic DNA"
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 17:
TGCAGGCAAC TTTGGCATGC 20

(2) INFORMATION FOR SEQ ID NO: 18:
(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 30 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear
(ii) MOLECULE TYPE: other nucleic acid
(A) DESCRIPTION: /desc = "Synthetic DNA"
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 18:
AGCAACACGA GCAAGATGAG GAGGATGACT 30

(2) INFORMATION FOR SEQ ID NO: 19:
(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 28 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear
(ii) MOLECULE TYPE: other nucleic acid
(A) DESCRIPTION: /desc = "Synthetic DNA"
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 19:
CCGGATCCTT CAACCCTTCT TCGATTTC 28

(2) INFORMATION FOR SEQ ID NO: 20:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 29 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear
(ii) MOLECULE TYPE: other nucleic acid
(A) DESCRIPTION: /desc = "Synthetic DNA"
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 20:
CCGGATCCAC GGCAATGCAT CTTGAAACC 29

(2) INFORMATION FOR SEQ ID NO: 21:
(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 27 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear
(ii) MOLECULE TYPE: other nucleic acid
(A) DESCRIPTION: /desc = "Synthetic DNA"
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 21:
GGTTAGCATA TTGTTTGTAG CATTGGG 27

(2) INFORMATION FOR SEQ ID NO: 22:
(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 27 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear
(ii) MOLECULE TYPE: other nucleic acid
(A) DESCRIPTION: /desc = "Synthetic DNA"
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 22:
ATCAATGAAA TATGTATAGT TCATAGC 27

(2) INFORMATION FOR SEQ ID NO: 23:
(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 27 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear
(ii) MOLECULE TYPE: other nucleic acid
(A) DESCRIPTION: /desc = "Synthetic DNA"
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 23:

CTTTCGTTCT TTTGGTTTTG CCATGGC 27

(2) INFORMATION FOR SEQ ID NO: 24:
(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 27 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear
(ii) MOLECULE TYPE: other nucleic acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 24:
AGACTTTTTA CAAACAAGAT AAATCCC 27

Claims

1. A DNA coding for any one of the following proteins (A) to (C):

   (A) a protein having a cellulose synthase activity and comprising an amino acid sequence shown in SEQ ID NO: 2 or an amino acid sequence involving deletion, substitution, insertion, or addition of one or several amino acids relevant to SEQ ID NO: 2;
   (B) a protein having a cellulose synthase activity and comprising an amino acid sequence shown in SEQ ID NO: 4 or an amino acid sequence involving deletion, substitution, insertion, or addition of one or several amino acids relevant to SEQ ID NO: 4; and
   (C) a protein having a cellulose synthase activity and comprising an amino acid sequence shown in SEQ ID NO: 8 or an amino acid sequence involving deletion, substitution, insertion, or addition of one or several amino acids relevant to SEQ ID NO: 8, and an amino acid sequence shown in SEQ ID NO: 11 or an amino acid sequence involving deletion, substitution, insertion, or addition of one or several amino acids relevant to SEQ ID NO: 11.

2. A recombinant vector comprising all or a part of the DNA as defined in claim 1.

3. A transformed cell transformed with the DNA as defined in claim 1.

4. A method for controlling cellulose synthesis in a cell, comprising the steps of introducing the DNA as defined in claim 1 into the cell, and expressing RNA having a nucleotide sequence homologous to the DNA as defined in claim 1 or a nucleotide sequence complementary to the DNA as defined in claim 1.

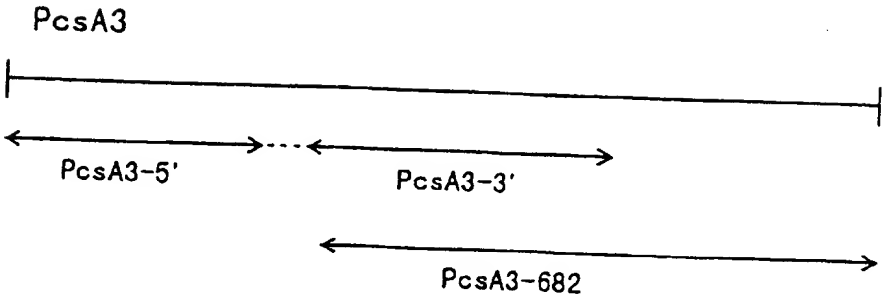

FIG. 1

SEQ ID NO: 14

```
5'  AATTCGGCACGAG  3'
3'        GCCGTGCTC  5' ---
```

FIG. 2

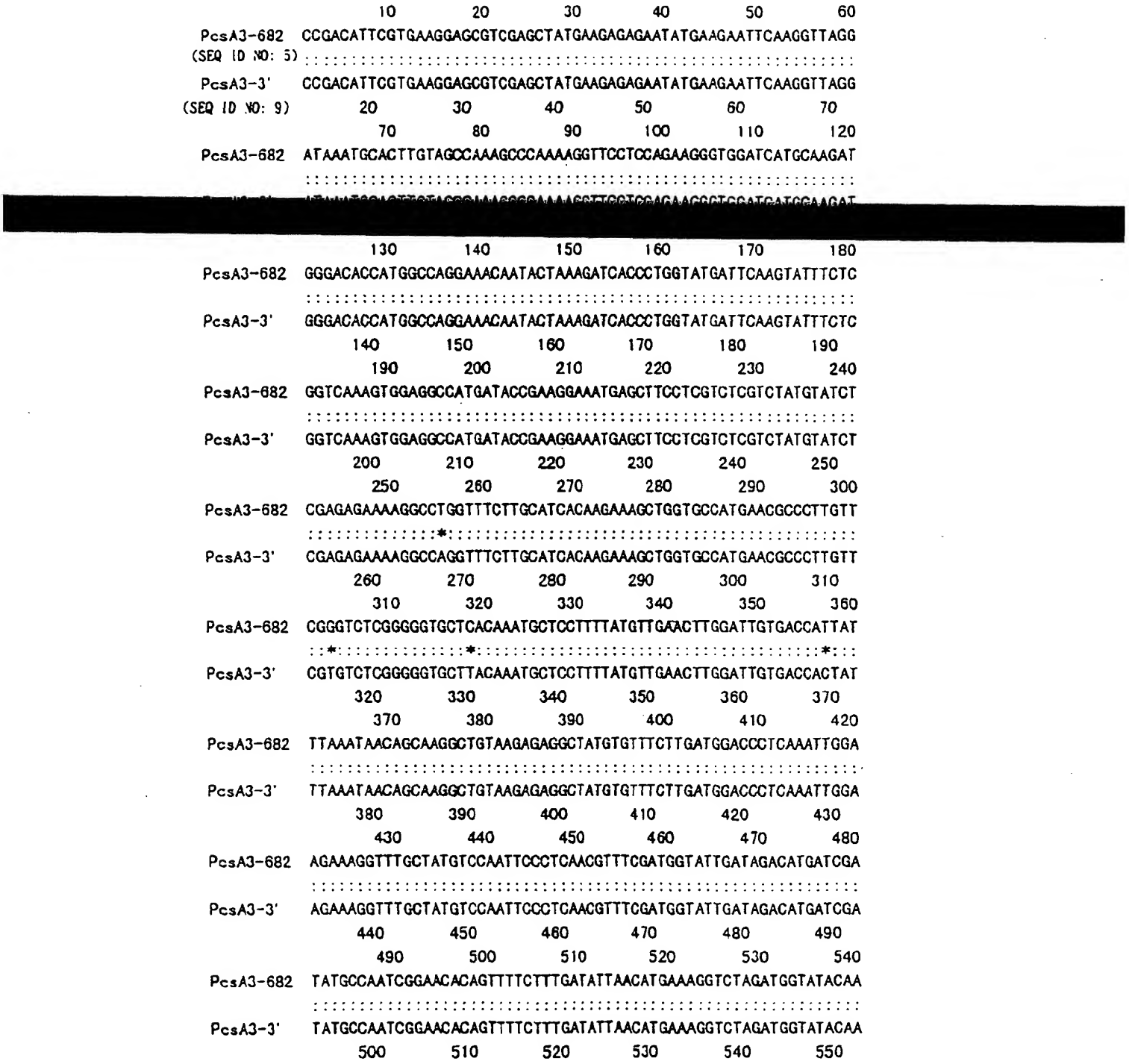

FIG. 3

```
                       550       560       570       580       590       600
PcsA3-682       GGCCCTGTATATGTCGGCACGGGGTGTGTTTTCAGAAGGCAAGCTCTTTATGGTTATGAA
(SEQ ID NO: 5)  ::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::
PcsA3-3'        GGCCCTGTATATGTCGGCACGGGGTGTGTTTTCAGAAGGCAAGCTCTTTATGGTTATGAA
(SEQ ID NO: 9)       560       570       580       590       600       610
                       610       620       630       640       650       660
PcsA3-682       CCTCCAAAGGGACCTAAGCGCCCGAAAATGGTAACCTGTGGTTGCTGCCCTTGTTTTGGA
                :::::::::::::::::::::::::::::::::::::::::::::::::::::*::::::
PcsA3-3'        CCTCCAAAGGGACCTAAGCGCCCGAAAATGGTAACCTGTGGTTGCTGCCCTTGCTTTGGA
                     620       630       640       650       660       670
                       670       680       690       700       710       720
PcsA3-682       CGCCGCAGAAAGGACAAAAAGCACTCTAAGGATGGTGGAAATGCAAATGGTCTAAGCCTA
                ::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::
PcsA3-3'        CGCCGCAGAAAGGACAAAAAGCACTCTAAGGATGGTGGAAATGCAAATGGTCTAAGCCTA
                     680       690       700       710       720       730
                       730       740       750       760       770       780
PcsA3-682       GAAGCAGCCAAAGATGACAAGGAGTTATTGATGTCCCACATGAACTTTGAAAAGAAATTT
                :::::::::*::::::::::::::::::::::::::::::::::::::::::::::::::
PcsA3-3'        GAAGCAGCCGAAGATGACAAGGAGTTATTGATGTCCCACATGAACTTTGAAAAGAAATTT
                     740       750       760       770       780       790
                       790       800       810       820       830       840
PcsA3-682       GGACAATCAGCCATTTTTGTAACTTCAACACTGATGGAACAAGGTGGTGTCCCTCCTTCT
                ::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::
PcsA3-3'        GGACAATCAGCCATTTTTGTAACTTCAACACTGATGGAACAAGGTGGTGTCCCTCCTTCT
                     800       810       820       830       840       850
                       850       860       870       880       890       900
PcsA3-682       TCAAGCCCCGCAGCTTTGCTCAAAGAAGCCATTCATGTAATTAGTTGTGGTTATGAAGAC
                ::::::::*:::::::::::::::::::::::::::::::::::::::::::::::::::
PcsA3-3'        TCAAGCCCTGCAGCTTTGCTCAAAGAAGCCATTCATGTAATTAGTTGTGGTTATGAAGAC
                     860       870       880       890       900       910
                       910       920       930       940       950       960
PcsA3-682       AAAACAGAATGGGGAAGCGAGCTTGGCTGGATTTACGGCTCGATTACAGAAGATATCTTA
                :::::*::::::::::::::::::::::::::::::::::::::::::::::::::::::
PcsA3-3'        AAAACCGAATGGGGAAGCGAGCTTGGCTGGATTTACGGCTCGATTACAGAAGATATCTTA
                     920       930       940       950       960       970
                       970       980
PcsA3-682       ACAGGATTCAAGATGCATTGCCGTGGAT
                :::::*::::::::::::::::::::::
PcsA3-3'        ACAGGTTTCAAGATGCATTGCCGTGGAT
                     980       990      1000
```

FIG. 4

[redacted]

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(54) Cellulose synthase gene

(57) mRNA was extracted at the stage for cotton plant fibrous cells to accumulate cellulose, and cDNA's complementary thereto were synthesized to construct a cDNA library. Clones of a number of 750 were arbitrarily selected from the library, and they were randomly subjected from to sequencing. Those having homology to an amino acid sequence deduced from a gene of cellulose 4-β-glucosyltransferase (bcsA) of cellulose synthase operon of acetic acid bacterium were selected from obtained nucleotide sequences of the respective clones. Thus, DNA coding for cellulose synthase was obtained.

EP 0 875 575 A3

European Patent Office

EUROPEAN SEARCH REPORT

Application Number
EP 98 30 2489

DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.6)
P,X	WO 98 00549 A (THE AUSTRALIAN NATIONAL [redacted] 8 January 1998 * page 1, line 3 - line 11 * * page 2, line 21 - page 7, line 28 * * example 8 * 'Sequence Listing: SEQ ID NO.9 and 10' ---	1-4	C12N15/54
X,D	PEAR, J.R. ETAL.: "Higher plants contain homologs of the bacterial celA genes encoding the catalytic subunit of cellulose synthase" PROC.NATL.ACAD.SCI.USA, vol. 93, October 1996, pages 12637-12642, XP002061424 * the whole document * ---	1-4	
Y	WO 91 13988 A (THE BOARD OF REGENTS, THE UNIVERSITY OF TEXAS SYSTEM) 19 September 1991 * page 1, line 18 - line 26 * * page 5, line 15 - page 8, line 13 * * figure 1; examples I,II,IV,V * ---	1-4	TECHNICAL FIELDS SEARCHED (Int.Cl.6) C12N
Y	LI, L. ET AL.: "ß-Glucan synthesis in the cotton fiber" PLANT PHYSIOLOGY, vol. 101, no. 4, 1993, pages 1149-1156, XP002087180 * page 1149 * * page 1154 - page 1155 * 'Abstract' and 'Discussion' ---	1-4	
E	WO 98 18949 A (CALGENE, INC.) 7 May 1998 * page 7, line 14 - page 9, line 25 * * figures 3,6,8; examples 1-7 * -----	1-4	

~~The present search report has been drawn up for all claims~~

Place of search	Date of completion of the search	Examiner
MUNICH	8 December 1998	Donath, C

CATEGORY OF CITED DOCUMENTS

X : particularly relevant if taken alone
Y : particularly relevant if combined with another document of the same category
A : technological background
O : non-written disclosure
P : intermediate document

T : theory or principle underlying the invention
E : earlier patent document, but published on, or after the filing date
D : document cited in the application
L : document cited for other reasons

& : member of the same patent family, corresponding document

EPO FORM 1503 03.82 (P04C01)

European Patent Office

Application Number
EP 98 30 2489

CLAIMS INCURRING FEES

The present European patent application comprised at the time of filing more than ten claims.

☐ Only part of the claims have been paid within the prescribed time limit. The present European search report has been drawn up for the first ten claims and for those claims for which claims fees have been paid, namely claim(s):

☐ No claims fees have been paid within the prescribed time limit. The present European search report has been drawn up for the first ten claims.

LACK OF UNITY OF INVENTION

The Search Division considers that the present European patent application does not comply with the requirements of unity of invention and relates to several inventions or groups of inventions, namely:

see sheet B

☐ All further search fees have been paid within the fixed time limit. The present European search report has been drawn up for all claims.

☐ As all searchable claims could be searched without effort justifying an additional fee, the Search Division did not invite payment of any additional fee.

☐ Only part of the further search fees have been paid within the fixed time limit. The present European search report has been drawn up for those parts of the European patent application which relate to the inventions in respect of which search fees have been paid, namely claims:

☒ None of the further search fees have been paid within the fixed time limit. The present European search report has been drawn up for those parts of the European patent application which relate to the invention first mentioned in the claims, namely claims:

1-4 (partially)

European Patent Office

LACK OF UNITY OF INVENTION
SHEET B

Application Number
EP 98 30 2489

The Search Division considers that the present European patent application does not comply with the requirements of unity of invention and relates to several inventions or groups of inventions, namely:

1. Claims: 1-4 (partially)

[redacted] acid sequence shown in SEQ ID NO:2 or an amino acid sequence involving deletion, substitution, insertion, or addition of one or several amino acids relevant to SEQ ID NO:2, a recombinant vector comprising all or part of said DNA, a cell being transformed with said DNA , and a method for controlling cellulose synthesis in a cell by the use of said DNA.

2. Claims: 1-4 (partially)

Claims 1- 4 (partially) refer to a DNA coding for a protein having a cellulose synthase activity and comprising an amino acid sequence shown in SEQ ID NO:4 or an amino acid sequence involving deletion, substitution, insertion, or addition of one or several amino acids relevant to SEQ ID NO:4, a recombinant vector comprising all or part of said DNA, a cell being transformed with said DNA, and a method for controlling cellulose synthesis in a cell by the use of said DNA.

3. Claims: 1-4 (partially)

Claims 1- 4 (partially) refer to a DNA coding for a protein having a cellulose synthase activity and comprising an amino acid sequence shown in SEQ ID NO:8 and in SEQ ID NO:11 or an amino acid sequence involving deletion, substitution, insertion, or addition of one or several amino acids relevant to SEQ ID NO:8 and/or SEQ ID NO:11, a recombinant vector comprising all or part of said DNA, a cell being transformed with said DNA, and a method for controlling cellulose synthesis in a cell by the use of said DNA.

SEQ ID NO: 6 for PcsA3.

PcsA1 is different from CelA1 reported by Pear et al. (Proceeding of National Academy of Science, USA (1996), 93, 12637-12642) in nucleotide sequence by 28 nucleotides. As a result, the former is different from the latter in amino acid sequence encoded thereby by 10 amino acid residues. In general, the sugar chain specificity and the substrate specificity of the sugar chain transferase are extremely changed by point mutation of the nucleotide of DNA (Yamamoto and Hakomori, The Journal of Biological Chemistry (1990) 265, 19257-19262). Therefore, it is unclear whether or not CelA1 codes for a protein having the cellulose synthase activity. Incidentally, the 48th Arg, the 56th Ser, the 81st Asn, the 104th Ala, the 110th Ser, the 247th Asp, the 376th Asp, the 386th Ser, the 409th Arg, and the 649th Ser in the amino acid sequence encoded by CelAl correspond to Gln, Ile, Ser, Thr, Pro, Asn, Glu, Pro, His, and Gly in PcsA1 respectively.

PcsA2 of the present invention contains the same sequence as that of CelA2 reported by Pear et al. However, CelA2 has an incomplete length, and it does not contain the entire coding region. CelA2 corresponds to nucleotide numbers of 1083 to 3311 in the nucleotide sequence of PcsA2 shown in SEQ ID NO: 3.

Any of the amino acid sequences shown in SEQ ID NOs: 2, 4, 6, 8, 10, and 11 is a novel sequence. All genes having nucleotide sequences coding for the amino acid sequences are included in the present invention.

The amino acid sequences described above may include deletion, substitution, insertion, and/or addition of one or more amino acid residues provided that the characteristic of the gene of the present invention is not substantially affected. The deletion, substitution, insertion, and/or addition of one or more amino acid residues as described above is obtainable by modifying the DNA's coding for the amino acid sequences shown in SEQ ID NOs: 2, 4, 6, 8, 10, and 11 randomly in accordance with the ordinary mutation treatment or intentionally in accordance with the site-directed mutagenesis method. As described above, in general, the sugar chain specificity and the substrate specificity of the sugar chain transferase are extremely changed by point mutation of the nucleotide of DNA. Therefore, DNA coding for a protein having the cellulose synthase activity is selected from the modified DNA's. The cellulose synthase activity can be measured, for example, by means of the method described by T. Hayashi: Measuring-β-glucan deposition in plant cell walls, in Modern Methods of Plant Analysis: Plant Fibers, eds. H. F. Linskens and J. F. Jackson, Springer-Verlag, 10: 138-160 (1989).

Those harboring proteins or genes partially different from the sequences shown in Sequence Listing may exist depending on, for example, the variety of cotton plant or natural mutation. However, such genes are also included in the gene of the present invention. Such a gene may be obtained as DNA which is hybridizable under the stringent condition with all or a part of the coding region of the nucleotide sequence shown in SEQ ID NO: 1, 3, 5, 7, or 9. The "stringent condition" referred to herein indicates a condition under which a so-called specific hybrid is formed, and non-specific hybrid is not formed. It is difficult to definitely express such a condition by using a numerical value. However, for example, the stringent condition is exemplified by a condition under which nucleic acids having high homology, for example, DNA's having homology of not less than 80 % undergo hybridization with each other, and nucleic acids having homology lower than the above do not undergo hybridization with each other.

<5> Utilization of gene of the present invention

The DNA of the present invention makes it possible to control the cellulose synthesis in prokaryotic cells such as acetobacterium and/or eukaryotic cells such as yeasts belonging to, for example, the genus Saccharomyces, cells of plant such as cotton plant, and cultured cells of mammals and the like.

Specifically, the cellulose synthesis in the cells as described above can be facilitated, for example, by connecting a promoter to an upstream region of the DNA of the present invention, inserting an obtained fragment into an appropriate vector to construct a recombinant vector, and introducing the vector into the cells. Alternatively, the cellulose synthesis in the cells can be suppressed by introducing an antisense gene of the DNA of the present invention into the cells.

The promoter and the vector may be selected from those ordinarily utilized to express heterogeneous genes, and the method ordinarily employed to express heterogeneous genes may be used as the transformation method. Specifically, in the case of yeast, it is possible to use a protein-expressing kit produced by Invitrogen, i.e., Pichia Expression Kit, and a vector pPIC9 contained in this kit. For example, COS7 cells may be used as mammalian cultured cells, and a vector CDM8 may be used therefor.

The present invention provides the DNA coding for cellulose synthase. The DNA provides a new method for controlling cellulose production by incorporating the DNA into prokaryotic cells and eukaryotic cells.

Brief Description of the Drawings

Fig. 1 shows a relationship between two clones of PcsA3 as an embodiment of the DNA of the present invention. Regions interposed between arrows indicate regions for which nucleotide sequences have been determined. A dotted line indicates a region for which no nucleotide sequence has been determined.

Fig. 2 shows a structure of EcoRI adapter.

Fig. 3 shows comparison between sequences of PcsA3-682 and PcsA3-3' (former half).

Fig. 4 shows comparison between sequences of PcsA3-682 and PcsA3-3' (latter half). ":" indicates coincident nucleotides, and "*" indicates non-coincident nucleotides.

Best Mode for Carrying Out the Invention

Examples of the present invention will be explained below.

<1> Preparation of total RNA from cotton plant

Cotton plant (Gossypium hirsutum L.) Coker 312 was used as a material. Fiber cells on 16 to 18 days post anthesis [redacted] while being frozen with liquid nitrogen. Powdered fiber was transferred to a [redacted] which 375 mg of DTT as a powder was added, followed by addition of 200 ml of XT buffer (obtained by adjusting 0.2 M sodium borate containing 30 mM EDTA and 1 % SDS to be pH 9.0, and then applying a diethylpyrocarbonate treatment, followed by autoclaving to obtain a solution to which vanadylribonucleoside was added to give a concentration of 10 mM) having been heated to 90 to 95 °C. An obtained solution was sufficiently agitated.

The solution was added with 100 mg of protease K, and it was agitated again. The solution was incubated at 40 °C for 2 hours, and then it was added with 16 ml of 2 M KCl. The solution was sufficiently agitated again, and it was left to stationarily stand in ice for 1 hour, followed by centrifugation for 20 minutes (4 °C) at 12,000 g by using a high speed refrigerated centrifuge.

An obtained supernatant was filtrated, and floating matters were removed. The solution was transferred to a measuring cylinder to measure the volume. The solution was transferred to another centrifuge tube, to which lithium chloride was added in an amount of 85 mg per 1 ml of the extract solution to give a final concentration of 2 M. The solution was left to stationarily stand at 4 °C overnight, and then precipitated RNA was separated by centrifugation for 20 minutes at 12,000 g. An obtained precipitate of RNA was washed and precipitated twice with cooled 2 M lithium chloride.

The obtained RNA was dissolved in 10 mM Tris buffer (pH 7.5) to give a concentration of about 2 mg/ml, to which 5 M potassium acetate was added to give a concentration of 200 mM. Ethanol was added thereto to give a concentration of 70 %, followed by cooling at -80 °C for 10 minutes. Centrifugation was performed at 4 °C for 10 minutes at 15,000 rpm, and then an obtained precipitate was suspended in an appropriate amount of sterilized water to give an RNA sample. As a result of quantitative measurement for the RNA sample, total RNA was obtained in an amount of 2 mg.

<2> Purification of mRNA

mRNA was purified as a poly(A)$^+$ RNA fraction from the total RNA obtained as described above. Purification was performed by using Oligotex-dT30 <Super> (purchased from Toyobo) as oligo(dT)-immobilized latex for poly(A)$^+$ RNA purification.

Elution buffer (10 mM Tris-HCl (pH 7.5), 1 mM EDTA, 0.1 % SDS) was added to a solution containing 1 mg of the total RNA to give a total volume of 1 ml, to which 1 ml of Oligotex-dT30 <Super> was added, followed by heating at 65 °C for 5 minutes and quick cooling on ice for 3 minutes. The obtained solution was added with 0.2 ml of 5 M NaCl, and it was incubated at 37 °C for 10 minutes, followed by centrifugation at 15,000 rpm for 3 minutes. After that, a supernatant was carefully removed.

An obtained pellet was suspended in 2.5 ml of Washing Buffer (10 mM Tris-HCl (pH 7.5), 1 mM EDTA, 0.5 M NaCl, 0.1 % SDS), and the suspension was centrifuged at 15,000 rpm for 3 minutes. After that, a supernatant was carefully removed. An obtained pellet was suspended in 1 ml of TE Buffer, and then it was heated at 65 °C 5 minutes. The suspension was quickly cooled on ice for 3 minutes, and then it was centrifuged at 15,000 rpm for 3 minutes to recover poly(A)$^+$ mRNA contained in an obtained supernatant.

Thus, the poly(A)$^+$ mRNA in an amount of about 10 μg was obtained from 1 mg of the total RNA. An aliquot of 5 μg thereof was used to prepare a cDNA library.

<3> Preparation of cDNA library

(1) Synthesis of cDNA

The mRNA obtained as described above was used as a template to synthesis cDNA by using a λZAP cDNA synthesis kit produced by Stratagene. The following solution was prepared and mixed in a tube.

5.0 µl 10 x 1st Strand Buffer (buffer for reverse transcription reaction);
3.0 µl 10 mM 1st Strand Methyl Nucleotide Mix (5-methyl dCTP, dATP, dGTP, dTTP mixture);
2.0 µl Linker-Primer (linker and primer);
H_2O (adjusted to give a total volume of 50 µl);
1.0 µl RNase Block II (RNase inhibitor).

The respective components described above were contents of the kit. Linker-Primer had a sequence as shown in SEQ ID NO: 13. Methylated nucleotide was used because it was intended not to allow cDNA to be digested by the restriction enzyme reaction performed later on. The reaction solution was agitated well, and then 5.0 µg of poly(A)$^+$ mRNA was added thereto, followed by being left to stand at room temperature for 10 minutes. Further, 2.5 µl of M-MuLV RTase (reverse transcriptase) was added (at this time, the total volume was 50 µl). The reaction solution was gently mixed, followed by centrifugation under a mild condition to allow the reaction solution to fall to the bottom of the tube. The reaction was performed at 37 °C for 60 minutes.

Next, the following solution was prepared and mixed in the tube in a certain order.

45.0 µl reaction solution containing cDNA primary chain;
40.0 µl 10 x 2nd Strand Buffer (buffer for polymerase reaction);
6.0 µl 2nd Strand Nucleotide Mixture (A, G, C, T mixture);
302.0 µl H_2O.

The following solution was further added. However, in order to allow RNase and DNA polymerase to simultaneously act, enzyme solutions were allowed to adhere to the wall of the tube. After that, a vortex treatment was promptly performed, and the reaction solutions were allowed to fall to the bottom of the tube by means of centrifugation to perform a reaction for synthesizing cDNA second strand at 16 °C for 150 minutes.

0.8 µl RNase H (RNA-degrading enzyme);
7.5 µl DNA polymerase I (10.0 u/µl).

The reaction solution was added with 400 µl of a mixed solution of phenol: chloroform (1:1). Agitation was performed well, followed by centrifugation at room temperature for 2 minutes. An obtained supernatant was added with 400 µl of phenol: chloroform again, which was subjected to a vortex treatment and centrifugation at room temperature for 2 minutes. An obtained supernatant was added with the following solution to precipitate cDNA.

33.3 µl 3 M sodium acetate solution;
867.0 µl 100 % ethanol.

The obtained solution was left to stand at -20 °C overnight, and it was centrifuged at room temperature for 60 minutes. After that, washing was gently performed with 80 % ethanol, followed by centrifugation for 2 minutes. A supernatant was removed. An obtained pellet was dried, and it was dissolved in 43.5 µl of sterilized water. An aliquot (39.0 µl) was added with the following solution to blunt-end cDNA terminals.

5.0 µl 10 x T4 DNA Polymerase Buffer (buffer for T4 polymerase reaction);
2.5 µl 2.5 mM dNTP Mix (A, G, C, T mixture);
3.5 µl T4 DNA polymerase (2.9 u/µl).

The reaction was performed at 37 °C for 30 minutes, to which 50 µl of distilled water was added, and then 100 µl of phenol: chloroform was added thereto, followed by a vortex treatment and centrifugation for 2 minutes. An obtained supernatant was added with 100 µl of chloroform, which was subjected to a vortex treatment, followed by centrifugation for 2 minutes. The supernatant was added with the following solution to precipitate cDNA.

7.0 µl 3 M sodium acetate solution;
226 µl 100 % ethanol.

The solution was left to stand on ice for 30 minutes or more, and it was centrifuged at 4 °C for 60 minutes. An obtained precipitate was washed with 150 µl of 80 % ethanol, followed by centrifugation for 2 minutes and drying. The cDNA pellet was dissolved in 7.0 µl of EcoRI Adaptor solution, to which the following solution was added to ligate the EcoRI adapter to both ends of the cDNA. Sequences of respective strands of the EcoRI adapter are shown in SEQ ID NO: 14 and Fig. 2.

1.0 µl 10 x Ligation Buffer (buffer for ligase reaction);
1.0 µl 10 mM ATP;
1.0 µl T4 DNA ligase.

The reaction solution was centrifuged under a mild condition, and it was left to stand at 4 °C overnight or more. The solution was treated at 70 °C for 30 minutes, and then it was centrifuged under a mild condition, followed by being left to stand at room temperature for 5 minutes. The reaction solution was added with the following solution to phosphorylate 5'-terminals of the EcoRI adapter.

1.0 µl 10 x Ligation Buffer (buffer for ligase reaction);
2.0 µl 10 mM ATP;

[redacted]

The reaction was performed at 37 °C for 30 minutes, followed by a treatment at 70 °C for 30 minutes. The solution was centrifuged under a mild condition, and it was left to stand at room temperature for 5 minutes. The following solution was further added thereto to perform a reaction at 37 °C for 90 minutes so that the XhoI site introduced by Linker-Primer was digested with XhoI, followed by being left to stand at room temperature to perform cooling.

28.0 µl XhoI Buffer;
3.0 µl XhoI (45 u/µl).

The reaction solution was added with 5.0 µl of 10 x STE (10 mM Tris-HCl (pH 8.0), 100 mM NaCl, 1 mM EDTA), which was added into a centrifuge column for removing short fragments (Sephacryl Spin Column) to perform centrifugation at 600 g for 2 minutes to obtain an eluent which was designated as Fraction 1. This operation was further repeated three times to obtain Fractions 2, 3, and 4 respectively. Fractions 3 and 4 were combined, to which phenol: chloroform (1:1) was added and agitated well, followed by centrifugation at room temperature for 2 minutes. An obtained supernatant was added with an equal amount of chloroform, and an obtained mixture was agitated well. The mixture was centrifuged at room temperature for 2 minutes to obtain a supernatant to which a two-fold amount of 100 % ethanol was added, followed by being left to stand at - 20 °C overnight. The solution was centrifuged at 4 °C for 60 minutes, followed by washing with an equal amount of 80 % ethanol. The solution was centrifuged at 4 °C for 60 minutes to obtain a cDNA pellet which was suspended in 10 µl of sterilized water.

(2) Preparation of cDNA library

The double strand cDNA obtained as described above was ligated with λ phage expression vector to prepare a recombinant vector. The following solution was prepared and mixed in a tube to perform a reaction at 12 °C overnight, followed by being left to stand at room temperature for 2 hours to ligate cDNA with the vector.

2.5 µl cDNA solution;
0.5 µl 10 x Ligation Buffer;
0.5 µl 10 mM ATP;
1.0 µl λZAP vector DNA (1 µg/µl);
0.5 µl T4 DNA ligase (4 Weiss u/µl).

(3) Packaging of phage DNA into phage particles

The phage vector containing the cDNA was packaged into phage particles by using an in vitro packaging kit (Gigapack II Gold packaging extract: produced by Stratagene). The recombinant phage solution was added to Freeze/Thaw extract immediately after dissolution, and the solution was placed on ice, to which 15 µl of Sonic extract was added to perform mixing well by pipetting. The reaction solution was centrifuged under a mild condition, and it was left to stand at room temperature (22 °C) for 2 hours. The reaction solution was added with 500 µl of Phage Dilution Buffer, to which 20 µl of chloroform was further added, followed by mixing. In order to measure the titer of the library, an aliquot (2 µl) of 500 µl of the aqueous phase was diluted in a ratio of 1:10 with 18 µl of SM buffer (5.8 g of NaCl, 2 g of $MgSo_4 \cdot 7H_2O$, 50 ml of 1 M Tris-HCl (pH 7.5), and 5 ml of 2 % gelatin in 1 L). The diluted solution (1 µl) and the phage stock solution (1 µl) were plated respectively together with 200 p1 of a culture solution of Escherichia coli PLK-F' strain having been cultivated to arrive at a value of OD_{600} of 0.5. That is, Escherichia coli PLK-F' strain was mixed with the phage solution to perform cultivation at 37 °C for 15 minutes. The obtained culture was added to 2 to 3 ml of top agar

(48 °C), which was immediately overlaid on NZY agar plate having been warmed at 37 °C. Cultivation was performed overnight at 37 °C. and appeared plaques were counted to calculate the titer. As a result, the titer was 1.2×10^6 pfu/ml.

(4) Amplification of library

A centrifuge tube was added with the packaging solution containing about 50.000 recombinant bacteriophages and 600 µl of a culture solution of Escherichia coli PLK-F' strain having been cultivated to have a value of OD_{600} of 0.5, followed by cultivation at 37 °C for 15 minutes. The culture solution was added with 6.5 ml of top agar having been maintained at 48 °C after dissolution, which was overlaid on 150 mm NZY plate having been warmed at about 37 °C, followed by cultivation at 37 °C for 5 to 8 hours. The respective plates were added with 10 ml of SM Buffer to perform cultivation at 4 °C overnight with gentle shaking. SM Buffer in the respective plates was collected in a sterilized polypropylene tube. The respective plates were rinsed with 2 ml of SM Buffer, and the rinsing solutions were collected in the same tube. Chloroform in an amount corresponding to 5 % of the total amount was added and mixed, followed by being left to stand at room temperature for 15 minutes. Bacterial cells were removed by centrifugation at 4,000 g for 5 minutes. An obtained supernatant was added with chloroform in an amount corresponding to 0.3 % of the total amount, and it was stored at 4 °C. The titer of the library amplified as described above was measured in the same manner as described above. As a result, the titer was 2.3×10^9 pfu/ml.

(5) Excision of plasmid from phage DNA

In vivo excision of the plasmid portion from the recombinant phage DNA was performed. The following solution was mixed in 50 ml of a conical tube to cause infection at 37 °C for 15 minutes:

culture solution of Escherichia coli XL1-Blue (OD_{600} = 0.1) 200 µl;
phage solution after amplification 200 µl (> 1×10^5 phage particles);
helper phage R408 1 µl (> 1×10^6 pfu/ml).

The mixed solution was added with 5 ml of 2 x YT medium to perform cultivation at 37 °C for 3 hours with shaking. A heat treatment was applied thereto at 70 °C for 20 minutes, followed by centrifugation at 4,000 g for 5 minutes. An obtained supernatant was decanted and transferred to a sterilized tube. Centrifugation was performed to obtain a supernatant which was diluted 100 times to obtain a solution. An aliquot (20 µl) of the solution was mixed with 200 µl of a culture solution of Escherichia coli XL1-Blue having been cultivated to obtain a value of OD_{600} of 1.0 to cause infection at 37 ° C for 15 minutes. Aliquots (1 to 100 µl) of the culture solution were plated on LB plates containing ampicillin, followed by cultivation at 37 °C overnight. Appeared colonies were randomly selected. Selected colonies were added with glycerol, and they were stored at -80 °C.

(6) Preparation of plasmid

Plasmids were prepared by using Magic Mini-prep kit produced by Promega. The culture fluid of Escherichia coli harboring the plasmid having been stored at -80 °C was inoculated into 5 ml of 2 x YT medium, followed by cultivation at 37 °C overnight. Centrifugation was performed for 5 minutes (4,000 rpm, 4 °C), and a supernatant was removed by decantation. An obtained bacterial cell pellet was added with 1 ml of TE buffer, followed by a vortex treatment. An obtained bacterial cell suspension was transferred to an Eppendorf tube, followed by centrifugation for 5 minutes (5,000 rpm, 4 ° C). A resultant supernatant was removed by decantation.

An obtained bacterial cell pellet was added with 300 µl of Cell Resuspension Solution, and it was sufficiently suspended therein. An obtained suspension was transferred to an Eppendorf tube. The suspension was agitated for 2 minutes with a mixer, to which 300 µl of Cell Lysis Solution was added, followed by agitation until the suspension became transparent. Neutralization Solution (300 µl) was added thereto, and agitation was performed by shaking with the hand, followed by centrifugation for 10 minutes (15,000 rpm).

Only an obtained supernatant was transferred to a new Eppendorf tube (1.5 ml). A suction tube was prepared, to which a cock, a miniature column and a syringe (injector) were connected in this order. A resin in an amount of 1 ml was charged into the syringe. The supernatant was poured into the syringe, and agitation was performed well, followed by suction. Column Washing Solution in an amount of 2 ml was added, and washing was performed while performing suction. Suction was continued for 1 to 2 minutes in order to dry up. The miniature column was removed from the equipment, and it was set in a new Eppendorf tube (1.5 ml). Sterilized water in an amount of 100 µl having been warmed at 65 to 70 °C was poured into the miniature column, and the column and the Eppendorf tube were centrifuged together for 1 minute (5,000 rpm). An eluted solution was transferred to an Eppendorf tube, to which 5 µl of 3 M sodium acetate aqueous solution was added, and 250 µl of cold ethanol was added thereto. The solution was centrifuged (15,000 rpm,

25 minutes), and a supernatant was discarded. An obtained precipitate was added with 1 ml of 70 % ethanol, followed by centrifugation again (15,000 rpm, 3 minutes). Ethanol was completely removed, and the tube was vacuum-dried in a desiccator. The precipitate was sufficiently dissolved in 20 µl of sterilized water, and an obtained solution was stored at -20 °C. An aliquot (1 µl) of the solution was dispensed, and it was subjected to electrophoresis together with volume markers to quantitatively determine the plasmid DNA.

<4> Determination of nucleotide sequence of cDNA and homology search with gene data base

(1) Determination of nucleotide sequence of cDNA

The nucleotide sequence of cDNA was analyzed by using DNA automatic sequencer 373A produced by Applied [redacted] was determined for about 750 clones which were randomly selected.

(2) Homology search

Partial sequences of about 750 clones were searched with a computer using BlastX. As a result, three clones appeared to be homologues of bacterial cellulose synthase subunit. Therefore, it was tried to isolate full length clones.

<5> Isolation of full length clones

(1) 5'-RACE

As a result of the homology search, the obtained homologue clones were found to be partial length clones. Therefore, primers were synthesized to make elongation toward the 5' upstream so that RT-PCR was performed by using mRNA as a template.

(1-a) Synthesis of first-strand DNA

The following solution was prepared and mixed in a tube.

0.5 µl 10 µmol gene-specific primer 1;
1 pg total RNA;
DEPC-treated H_2O (adjusted to give a total amount of 9 µl).

The following oligonucleotides were used as the gene-specific primer 1. That is, an oligonucleotide having a nucleotide sequence shown in SEQ ID NO: 15 was used for PcsA1. An oligonucleotide having a nucleotide sequence shown in SEQ ID NO: 16 was used for PcsA2. An oligonucleotide having a nucleotide sequence shown in SEQ ID NO: 17 was used for PcsA3.

The reaction solution was gently mixed, and then it was centrifuged under a mild condition to allow the reaction solution to fall to the bottom of the tube. The solution was left to stand at 70 °C for 10 minutes, followed by immediate cooling on ice.

Next, the following solution was prepared and mixed in the tube.

5 x RT Buffer 5 µl;
25 mM $MgCl_2$ 2.5 µl;
2 mM dNTP mix 5 µl;
0.1 M DTT 2.5 µl;
H_2O (added to give a total amount of 24 µl).

The solution was gently agitated, and then it was centrifuged under a mild condition to allow the reaction solution to fall to the bottom of the tube, followed by being left to stand at 42 °C for 1 minute. The solution was added with 1 µl of SuperScriptII RT (reverse transcriptase, GIBCO BRL), and it was gently mixed. After that, the reaction was performed at 42 °C for 50 minutes. Subsequently, the reaction solution was left to stand at 70 °C for 15 minutes to stop the reaction. Centrifugation was performed under a mild condition to allow the reaction solution to fall to the bottom of the tube, followed by being left to stand at 37 °C. RNase H (produced by Toyobo) in an amount of 1 µl was added thereto to perform a reaction at 37 °C for 30 minutes.

Subsequently. in order to remove excessive primers and nucleotides contained in the reaction solution, gel filtration was performed by using a purification column produced by Boehringer, Quick Spin Columns. At first, the tip of the column was removed. followed by centrifugation at 1,100 x g for 2 minutes to discard the buffer. The reaction solution was introduced into the central area of the column, followed by centrifugation at 1,100 x g for 4 minutes to recover the solution.

(1-b) Poly(dC) tailing

An aliquot (5 μl) was dispensed from the obtained solution, to which the following solution was added.

5 μl 5 x $CoCl_2$ Buffer;
2.5 μl 2 mM dCTP;
H_2O (adjusted to give a total amount of 24 μl).

The reaction solution was mixed well, and it was left to stand at 94 °C for 3 minutes. Centrifugation was performed under a mild condition to allow the reaction solution to fall to the bottom of the tube, followed by being left to stand on ice. Terminal transferase TdT (produced by Toyobo) was added thereto in an amount of 1 μl, followed by mixing under a mild condition to perform a reaction at 37 °C for 10 minutes. Subsequently, the reaction solution was left to stand at 65 °C for 10 minutes to stop the reaction.

(1-c) PCR reaction

An aliquot (2.5 μl) was dispensed from the reaction solution, to which the following solution was added.

2.5 μl 10 x PCR Buffer;
2.5 μl 2 mM dNTP mix;
0.5 μl Gene-specific primer 2;
0.5 μl Abridged Anchor Primer (GIBCO BRL);
0.5 μl Advantage Klentaq Polymerase Mix (Clontech);
H_2O (adjusted to give a total amount of 25 μl).

The following oligonucleotides were used as Gene-specific primer 2. That is, an oligonucleotide having a nucleotide sequence shown in SEQ ID NO: 18 was used for PcsA1. An oligonucleotide having a nucleotide sequence shown in SEQ ID NO: 19 was used for PcsA2. An oligonucleotide having a nucleotide sequence shown in SEQ ID NO: 20 was used for PcsA3.

The solution was introduced into a 0.2 ml tube to perform the PCR reaction under the following condition.

PAD	94 °C	90 seconds
30 cycles	94 °C	30 seconds
	60 to 68 °C	30 to 60 seconds
	68 °C	180 seconds
Final	68 °C	7 minutes
Hold	4 °C	

The reaction solution was subjected to agarose gel electrophoresis to extract, from the gel, DNA's corresponding to portions having the largest size (about 1.8 K for PcsA1, about 2 K for PcsA2, and about 2.2 K for PcsA3). GENO-BIND produced by CLONTECH was used for the extraction, and the procedure was carried out in accordance with its protocol. The DNA thus obtained was subjected to Poly(dC)tailing, which was used as a template to perform the PCR reaction. The condition and the composition of the reaction solution were the same as those described above.

(2) Cloning

(2-a) 5'-RACE TA cloning

Starting from the obtained PCR reaction solution, cloning was performed by using TA Cloning Kit produced by Invitrogen in accordance with its protocol.

The following solution was added to an aliquot (1.5 μl) of the PCR reaction solution obtained as described above.

0.5 µl 10 x Ligation Buffer;
1 µl pCRII vector;
0.5 µl T4 DNA Ligase;
1.5 µl dH_2O.

The reaction was performed at 14 °C overnight. An aliquot (2 p1) of the reaction solution was added to 25 p1 of Escherichia coli competent cell (JM109) preparation, followed by being left to stand for 30 minutes on ice. After that, heat shock was applied at 42 °C for 30 seconds. The solution was stationarily left to stand on ice for 2 minutes, to which 450 µl of SOB medium was thereafter added to perform cultivation at 37 °C for 1 hour with shaking at 200 rpm. The culture was spread over Amp/Xgal/IPTG plate, followed by incubation at 37 °C overnight. The plasmid was extracted from obtained colonies in accordance with the method as described above.

The procedure was carried out by using DNA Sequencer 377 produced by ABI in accordance with its protocol. The sequencing reaction was performed by using M13 primer and synthetic oligomer as primers, based on the use of Dye Terminater Cycle Sequencing Kit produced by the same company. As a result of the sequencing, as for PcsA3, it was revealed that another clone also belonging to the group of PcsA3 but having a slightly different sequence (one position for amino acid) was isolated (see Figs. 3 and 4). A nucleotide sequence of a clone (PcsA3-682) containing the 3'-side region of PcsA3 and an amino acid sequence deduced from this nucleotide sequence are shown in SEQ ID NOs: 5 and 6. A nucleotide sequence of a 5'-portion (PcsA3-5') of another clone containing the 5'-side region of PcsA3 and an amino acid sequence deduced from this nucleotide sequence are shown in SEQ ID NOs: 7 and 8. A nucleotide sequence of a 3'-portion (PcsA3-3') of the clone and an amino acid sequence deduced from this nucleotide sequence are shown in SEQ ID NOs: 9 and 10.

As for PcsAl and PcsA2, primers for 5'-terminal and 3'-terminal of a region containing ORF were synthesized on the basis of the obtained sequences to perform the PCR reaction. Thus, complete length clones were isolated by means of TA cloning. The condition and the composition of the reaction solution were the same as those described above.

Oligonucleotides shown in SEQ ID NO: 21 (5'-terminal) and SEQ ID NO: 22 (3'-terminal) were used as the primers for PcsAl. Oligonucleotides shown in SEQ ID NO: 23 (5'-terminal) and SEQ ID NO: 24 (3'-terminal) were used as the primers for PcsA2. Results are shown in SEQ ID NOs: 1 to 4.

Ann x to the description

SEQUENCE LISTING

(1) GENERAL INFORMATION:
(i) APPLICANT: NISSHINBO INDUSTRIES, INC.
HAYASHI, Takahisa
(ii) TITLE OF INVENTION: CELLULOSE SYNTHASE GENE
(iii) NUMBER OF SEQUENCES: 24
(iv) CORRESPONDENCE ADDRESS:
(A) ADDRESSEE:
(B) STREET:
(C) CITY:
(E) COUNTRY:
(F) ZIP:
(v) COMPUTER READABLE FORM:
(A) MEDIUM TYPE: Floppy disk
(B) COMPUTER: IBM PC compatible
(C) OPERATING SYSTEM: PC-DOS/MS-DOS
(D) SOFTWARE: PatentIn Release #1.0, Version #1.30
(vi) CURRENT APPLICATION DATA:
(A) APPLICATION NUMBER:
(B) FILING DATE:
(C) CLASSIFICATION:
(vii) PRIOR APPLICATION DATA:
(A) APPLICATION NUMBER: JP 9-83133
(B) FILING DATE: 1-APR-1997
(viii) ATTORNEY/AGENT INFORMATION:
(A) NAME:
(B) REGISTRATION NUMBER:
(ix) TELECOMMUNICATION INFORMATION:
(A) TELEPHONE:
(B) TELEFAX:

(2) INFORMATION FOR SEQ ID NO: 1:
(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 3207 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear
(ii) MOLECULE TYPE: cDNA to mRNA
(vi) ORIGINAL SOURCE:
(A) ORGANISM: Gossypium hirsutum L.

(C) INDIVIDUAL ISOLATE: Coker312

(ix) FEATURE:

(A) NAME/KEY: CDS

(B) LOCATION:77..3001

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 1:

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                    1               5                  10

[redacted]

Cys Gly Glu His Val Gly Leu Asn Val Asn Gly Glu Pro Phe Val Ala
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TGC CAT GAA TGT AAT TTC CCT ATT TGT AAG AGT TGT TTT GAG TAT GAT      205
Cys His Glu Cys Asn Phe Pro Ile Cys Lys Ser Cys Phe Glu Tyr Asp
            30                  35                  40
CTT AAG GAA GGA CAA AAA GCT TGC TTG CGT TGT GGT ATT CCG TAT GAT      253
Leu Lys Glu Gly Gln Lys Ala Cys Leu Arg Cys Gly Ile Pro Tyr Asp
        45                  50                  55
GAA AAC CTG TTG GAC GAT GTC GAG AAG GCC ACC GGC GAT CAA TCG ACA      301
Glu Asn Leu Leu Asp Asp Val Glu Lys Ala Thr Gly Asp Gln Ser Thr
    60                  65                  70                  75
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                80                  85                  90
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        110                 115                 120
AAC AAG AAG AAG AAG CCT GCA ACA ACT AAG GTT GAA AGA GAG GCT GAA      493
Asn Lys Lys Lys Lys Pro Ala Thr Thr Lys Val Glu Arg Glu Ala Glu
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ATC CCA CCT GAG CAA CAA ATG GAA GAT AAA CCG GCA CCG GAT GCT TCC      541
Ile Pro Pro Glu Gln Gln Met Glu Asp Lys Pro Ala Pro Asp Ala Ser
140                 145                 150                 155
CAG CCC CTC TCG ACT ATA ATT CCA ATC CCG AAA AGC AGA CTT GCA CCA      589
Gln Pro Leu Ser Thr Ile Ile Pro Ile Pro Lys Ser Arg Leu Ala Pro
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ACT TCA GTC ATA TGT GAA ATC TGG TTT GCT TTT TCC TGG GTG TTG GAT      733
Thr Ser Val Ile Cys Glu Ile Trp Phe Ala Phe Ser Trp Val Leu Asp
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CAG TTC CCT AAG TGG TAT CCT GTT AAC AGG GAA ACA TAC ATT GAC AGA      781
Gln Phe Pro Lys Trp Tyr Pro Val Asn Arg Glu Thr Tyr Ile Asp Arg
220                 225                 230                 235
CTG TCT GCA AGA TAT GAA AGA GAA GGT GAA CCT AAT GAA CTT GCT GCA      829
Leu Ser Ala Arg Tyr Glu Arg Glu Gly Glu Pro Asn Glu Leu Ala Ala
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GTT GAC TTC TTT GTG AGT ACA GTG GAT CCA TTG AAA GAG CCT CCA TTG      877
Val Asp Phe Phe Val Ser Thr Val Asp Pro Leu Lys Glu Pro Pro Leu
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ATT ACT GCC AAT ACT GTG CTT TCC ATC CTT GCC TTG GAC TAC CCG GTA      925
Ile Thr Ala Asn Thr Val Leu Ser Ile Leu Ala Leu Asp Tyr Pro Val
        270                 275                 280
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Asp Lys Val Ser Cys Tyr Ile Ser Asp Asp Gly Ala Ala Met Leu Thr
    285                 290                 295
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300                 305                 310                 315
TTC TGC AAA AAA TTT TCC ATT GAA CCA CGG GCA CCT GAG TTT TAC TTC     1069
Phe Cys Lys Lys Phe Ser Ile Glu Pro Arg Ala Pro Glu Phe Tyr Phe
                320                 325                 330
TCA CAG AAG ATT GAT TAC TTG AAA GAT AAA GTG CAG CCC TCT TTT GTA     1117
Ser Gln Lys Ile Asp Tyr Leu Lys Asp Lys Val Gln Pro Ser Phe Val
            335                 340                 345
AAA GAA CGT AGA GCT ATG AAA AGA GAT TAC GAA GAG TAC AAA ATT CGA     1165
Lys Glu Arg Arg Ala Met Lys Arg Asp Tyr Glu Glu Tyr Lys Ile Arg
        350                 355                 360
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ACA ATG CAA GAT GGA ACT CCT TGG CCG GGA AAT AAC CCG CGT GAT CAC     1261
Thr Met Gln Asp Gly Thr Pro Trp Pro Gly Asn Asn Pro Arg Asp His
380                 385                 390                 395
CCT GGC ATG ATT CAG GTT TTC CTT GGA TAT AGC GGT GCT CAT GAC ATC     1309
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            415                 420                 425
CCT GGC TAC CAA CAC CAC AAA AAG GCT GGT GCT GAA AAT GCT TTG GTT   1405
Pro Gly Tyr Gln His His Lys Lys Ala Gly Ala Glu Asn Ala Leu Val
        430                 435                 440
AGG GTG TCT GCA GTT CTT ACA AAT GCT CCC TTC ATC CTC AAT CTT GAT   1453
[redacted]
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Cys Asp His Tyr Val Asn Asn Ser Lys Ala Val Arg Glu Ala Met Cys
460                 465                 470                 475
TTC TTG ATG GAC CCA CAA GTC GGT CGA GAT GTC TGC TAT GTG CAG TTT   1549
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                480                 485                 490
CCT CAA AGA TTT GAT GGC ATA GAT AGG AGT GAT CGA TAT GCC AAT CGG   1597
Pro Gln Arg Phe Asp Gly Ile Asp Arg Ser Asp Arg Tyr Ala Asn Arg
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AAC ACA GTT TTC TTT GAT GTT AAC ATG AAA GGT CTT GAT GGA ATC CAA   1645
Asn Thr Val Phe Phe Asp Val Asn Met Lys Gly Leu Asp Gly Ile Gln
        510                 515                 520
GGG CCT GTT TAT GTG GGA ACA GGT TGT GTT TTC AAT AGG CAA GCA CTT   1693
Gly Pro Val Tyr Val Gly Thr Gly Cys Val Phe Asn Arg Gln Ala Leu
    525                 530                 535
TAT GGC TAT GGT CCA CCT TCA ATG CCA AGT TTT CCC AAG TCA TCC TCC   1741
Tyr Gly Tyr Gly Pro Pro Ser Met Pro Ser Phe Pro Lys Ser Ser Ser
540                 545                 550                 555
TCA TCT TGC TCG TGT TGC TGC CCC GGC AAG AAG GAA CCT AAA GAT CCA   1789
Ser Ser Cys Ser Cys Cys Cys Pro Gly Lys Lys Glu Pro Lys Asp Pro
                560                 565                 570
TCA GAG CTT TAT AGG GAT GCA AAA CGG GAA GAA CTT GAT GCT GCC ATC   1837
Ser Glu Leu Tyr Arg Asp Ala Lys Arg Glu Glu Leu Asp Ala Ala Ile
            575                 580                 585
TTT AAC CTT AGG GAA ATT GAC AAT TAT GAT GAG TAT GAA AGA TCA ATG   1885
Phe Asn Leu Arg Glu Ile Asp Asn Tyr Asp Glu Tyr Glu Arg Ser Met
        590                 595                 600
TTG ATC TCT CAA ACA AGC TTT GAG AAA ACT TTT GGC TTA TCT TCA GTC   1933
Leu Ile Ser Gln Thr Ser Phe Glu Lys Thr Phe Gly Leu Ser Ser Val
    605                 610                 615
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620                 625                 630                 635
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Asn Pro Ser Thr Leu Ile Lys Glu Ala Ile His Val Ile Gly Cys Gly
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TCA GTC ACT GAG GAT ATC TTA ACC GGC TTC AAA ATG CAC TGC CGA GGA      2125
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TGG AGA TCG ATT TAC TGC ATG CCC TTA AGG CCA GCA TTC AAA GGA TCT      2173
Trp Arg Ser Ile Tyr Cys Met Pro Leu Arg Pro Ala Phe Lys Gly Ser
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GCA CCC ATC AAT CTG TCT GAT CGG TTG CAC CAG GTT CTT CGA TGG GCT      2221
Ala Pro Ile Asn Leu Ser Asp Arg Leu His Gln Val Leu Arg Trp Ala
700                 705                 710                 715
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Leu Gly Ser Val Glu Ile Phe Leu Ser Arg His Cys Pro Leu Trp Tyr
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            735                 740                 745
AAC ACC ATT GTC TAT CCT TTC ACA TCC CTT CCA CTC ATT GCC TAT TGT      2365
Asn Thr Ile Val Tyr Pro Phe Thr Ser Leu Pro Leu Ile Ala Tyr Cys
        750                 755                 760
TCA CTA CCA GCA ATC TGT CTT CTC ACA GGA AAA TTT ATC ATA CCA ACG      2413
Ser Leu Pro Ala Ile Cys Leu Leu Thr Gly Lys Phe Ile Ile Pro Thr
    765                 770                 775
CTC TCA AAC CTG GCA AGT GTT CTC TTT CTT GGC CTT TTC CTT TCC ATT      2461
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780                 785                 790                 795
ATC GTG ACT GCT GTT CTC GAG CTC CGA TGG AGT GGT GTC AGC ATT GAG      2509
Ile Val Thr Ala Val Leu Glu Leu Arg Trp Ser Gly Val Ser Ile Glu
                800                 805                 810
GAC TTA TGG CGT AAC GAG CAG TTT TGG GTC ATC GGT GGC GTT TCA GCC      2557
Asp Leu Trp Arg Asn Glu Gln Phe Trp Val Ile Gly Gly Val Ser Ala
            815                 820                 825
CAT CTC TTT GCC GTC TTC CAA GGT TTC CTT AAG ATG CTT GCG GGC ATT      2605
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          830                 835                 840
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    845                 850                 855
GGT GAG CTC TAC ATT GTG AAA TGG ACT ACA CTT CTA ATC CCT CCA ACA      2701
Gly Glu Leu Tyr Ile Val Lys Trp Thr Thr Leu Leu Ile Pro Pro Thr
860                 865                 870                 875
ACA CTC CTC ATC GTC AAC ATG GTT GGT GTC GTT GCC GGA TTC TCC GAT      2749
Thr Leu Leu Ile Val Asn Met Val Gly Val Val Ala Gly Phe Ser Asp
[redacted]
GCC CTC AAC AAA GGG TAC GAA GCT TGG GGA CCA CTC TTT GGC AAA GTG      2797
Ala Leu Asn Lys Gly Tyr Glu Ala Trp Gly Pro Leu Phe Gly Lys Val
          895                 900                 905
TTC TTT TCC TTC TGG GTC ATC CTC CAT CTT TAT CCA TTC CTC AAA GGT      2845
Phe Phe Ser Phe Trp Val Ile Leu His Leu Tyr Pro Phe Leu Lys Gly
    910                 915                 920
CTT ATG GGA CGC CAA AAC AGG ACA CCA ACC ATT GTT GTC CTT TGG TCA      2893
Leu Met Gly Arg Gln Asn Arg Thr Pro Thr Ile Val Val Leu Trp Ser
925                 930                 935
GTG TTG TTG GCT TCT GTC TTC TCT CTT GTT TGG GTT CGG ATC AAC CCG      2941
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940                 945                 950                 955
TTT GTC AGC ACC GCC GAT AGC ACC ACC GTG TCA CAG AGC TGC ATT TCC      2989
Phe Val Ser Thr Ala Asp Ser Thr Thr Val Ser Gln Ser Cys Ile Ser
                960                 965                 970
ATT GAT TGT TGATGATATT ATGTGTTTCT TAGAATTGAA ATCATTGCAA              3038
Ile Asp Cys
GTAAGTGGAC TGAAACATGT CTATTGACTA AGTTTTGAAC AGTTTGTACC CATTTTATTC    3098
TTAGCAGTGT GTAATTTTCC TAAACAATGC TATGAACTAT ACATATTTCA TTGATATTTA    3158
CATTAAATGA AACTACATCA GTCTGCAGAA AAAAAAAAAA AAAAAAAAA                3207
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(2) INFORMATION FOR SEQ ID NO: 2:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 974 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 2:

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20 25 30

Phe Pro Ile Cys Lys Ser Cys Phe Glu Tyr Asp Leu Lys Glu Gly Gln

35 40 45

Lys Ala Cys Leu Arg Cys Gly Ile Pro Tyr Asp Glu Asn Leu Leu Asp

50 55 60

Asp Val Glu Lys Ala Thr Gly Asp Gln Ser Thr Met Ala Ala His Leu

65 70 75 80

Ser Lys Ser Gln Asp Val Gly Ile His Ala Arg His Ile Ser Ser Val

85 90 95

Ser Thr Leu Asp Ser Glu Met Thr Glu Asp Asn Gly Asn Pro Ile Trp

100 105 110

Lys Asn Arg Val Glu Ser Trp Lys Glu Lys Lys Asn Lys Lys Lys Lys

115 120 125

Pro Ala Thr Thr Lys Val Glu Arg Glu Ala Glu Ile Pro Pro Glu Gln

130 135 140

Gln Met Glu Asp Lys Pro Ala Pro Asp Ala Ser Gln Pro Leu Ser Thr

145 150 155 160

Ile Ile Pro Ile Pro Lys Ser Arg Leu Ala Pro Tyr Arg Thr Val Ile

165 170 175

Ile Met Arg Leu Ile Ile Leu Gly Leu Phe Phe His Tyr Arg Val Thr

180 185 190

Asn Pro Val Asp Ser Ala Phe Gly Leu Trp Leu Thr Ser Val Ile Cys

195 200 205

Glu Ile Trp Phe Ala Phe Ser Trp Val Leu Asp Gln Phe Pro Lys Trp

210 215 220

Tyr Pro Val Asn Arg Glu Thr Tyr Ile Asp Arg Leu Ser Ala Arg Tyr

225 230 235 240

Glu Arg Glu Gly Glu Pro Asn Glu Leu Ala Ala Val Asp Phe Phe Val

245 250 255

Ser Thr Val Asp Pro Leu Lys Glu Pro Pro Leu Ile Thr Ala Asn Thr

260 265 270

Val Leu Ser Ile Leu Ala Leu Asp Tyr Pro Val Asp Lys Val Ser Cys

275 280 285

Tyr Ile Ser Asp Asp Gly Ala Ala Met Leu Thr Phe Glu Ser Leu Val

290 295 300

Glu Thr Ala Asp Phe Ala Arg Lys Trp Val Pro Phe Cys Lys Lys Phe

305 310 315 320

Ser Ile Glu Pro Arg Ala Pro Glu Phe Tyr Phe Ser Gln Lys Ile Asp

325 330 335

Tyr Leu Lys Asp Lys Val Gln Pro Ser Phe Val Lys Glu Arg Arg Ala

340 345 350

Met Lys Arg Asp Tyr Glu Glu Tyr Lys Ile Arg Ile Asn Ala Leu Val
355 360 365
Ala Lys Ala Gln Lys Thr Pro Glu Glu Gly Trp Thr Met Gln Asp Gly
370 375 380
Thr Pro Trp Pro Gly Asn Asn Pro Arg Asp His Pro Gly Met Ile Gln
385 390 395 400
Val Phe Leu Gly Tyr Ser Gly Ala His Asp Ile Glu Gly Asn Glu Leu
405 410 415

[redacted]

His Lys Lys Ala Gly Ala Glu Asn Ala Leu Val Arg Val Ser Ala Val
435 440 445
Leu Thr Asn Ala Pro Phe Ile Leu Asn Leu Asp Cys Asp His Tyr Val
450 455 460
Asn Asn Ser Lys Ala Val Arg Glu Ala Met Cys Phe Leu Met Asp Pro
465 470 475 480
Gln Val Gly Arg Asp Val Cys Tyr Val Gln Phe Pro Gln Arg Phe Asp
485 490 495
Gly Ile Asp Arg Ser Asp Arg Tyr Ala Asn Arg Asn Thr Val Phe Phe
500 505 510
Asp Val Asn Met Lys Gly Leu Asp Gly Ile Gln Gly Pro Val Tyr Val
515 520 525
Gly Thr Gly Cys Val Phe Asn Arg Gln Ala Leu Tyr Gly Tyr Gly Pro
530 535 540
Pro Ser Met Pro Ser Phe Pro Lys Ser Ser Ser Ser Ser Cys Ser Cys
545 550 555 560
Cys Cys Pro Gly Lys Lys Glu Pro Lys Asp Pro Ser Glu Leu Tyr Arg
565 570 575
Asp Ala Lys Arg Glu Glu Leu Asp Ala Ala Ile Phe Asn Leu Arg Glu
580 585 590
Ile Asp Asn Tyr Asp Glu Tyr Glu Arg Ser Met Leu Ile Ser Gln Thr
595 600 605
Ser Phe Glu Lys Thr Phe Gly Leu Ser Ser Val Phe Ile Glu Ser Thr
610 615 620
Leu Met Glu Asn Gly Gly Val Ala Glu Ser Ala Asn Pro Ser Thr Leu
625 630 635 640
Ile Lys Glu Ala Ile His Val Ile Gly Cys Gly Tyr Glu Glu Lys Thr
645 650 655
Ala Trp Gly Lys Glu Ile Gly Trp Ile Tyr Gly Ser Val Thr Glu Asp
660 665 670
Ile Leu Thr Gly Phe Lys Met His Cys Arg Gly Trp Arg Ser Ile Tyr

675 680 685
Cys Met Pro Leu Arg Pro Ala Phe Lys Gly Ser Ala Pro Ile Asn Leu
690 695 700
Ser Asp Arg Leu His Gln Val Leu Arg Trp Ala Leu Gly Ser Val Glu
705 710 715 720
Ile Phe Leu Ser Arg His Cys Pro Leu Trp Tyr Gly Phe Gly Gly Gly
725 730 735
Arg Leu Lys Trp Leu Gln Arg Leu Ala Tyr Ile Asn Thr Ile Val Tyr
740 745 750
Pro Phe Thr Ser Leu Pro Leu Ile Ala Tyr Cys Ser Leu Pro Ala Ile
755 760 765
Cys Leu Leu Thr Gly Lys Phe Ile Ile Pro Thr Leu Ser Asn Leu Ala
770 775 780
Ser Val Leu Phe Leu Gly Leu Phe Leu Ser Ile Ile Val Thr Ala Val
785 790 795 800
Leu Glu Leu Arg Trp Ser Gly Val Ser Ile Glu Asp Leu Trp Arg Asn
805 810 815
Glu Gln Phe Trp Val Ile Gly Gly Val Ser Ala His Leu Phe Ala Val
820 825 830
Phe Gln Gly Phe Leu Lys Met Leu Ala Gly Ile Asp Thr Asn Phe Thr
835 840 845
Val Thr Ala Lys Ala Ala Asp Asp Ala Asp Phe Gly Glu Leu Tyr Ile
850 855 860
Val Lys Trp Thr Thr Leu Leu Ile Pro Pro Thr Thr Leu Leu Ile Val
865 870 875 880
Asn Met Val Gly Val Val Ala Gly Phe Ser Asp Ala Leu Asn Lys Gly
885 890 895
Tyr Glu Ala Trp Gly Pro Leu Phe Gly Lys Val Phe Phe Ser Phe Trp
900 905 910
Val Ile Leu His Leu Tyr Pro Phe Leu Lys Gly Leu Met Gly Arg Gln
915 920 925
Asn Arg Thr Pro Thr Ile Val Val Leu Trp Ser Val Leu Leu Ala Ser
930 935 940
Val Phe Ser Leu Val Trp Val Arg Ile Asn Pro Phe Val Ser Thr Ala
945 950 955 960
Asp Ser Thr Thr Val Ser Gln Ser Cys Ile Ser Ile Asp Cys
965 970

(2) INFORMATION FOR SEQ ID NO: 3:
(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 3311 base pairs

(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA to mRNA

(vi) ORIGINAL SOURCE:
(A) ORGANISM: Gossypium hirsutum L.
(C) INDIVIDUAL ISOLATE: Coker312

(ix) FEATURE:
(A) NAME/KEY: CDS

[redacted]

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 3:

```
CTTTCGTTCT TTTGGTTTTG CC ATG GCT TCA ACC ACC ATG GCC GCT GGC TTT        52
                         Met Ala Ser Thr Thr Met Ala Ala Gly Phe
                           1               5                  10
GGT TCA CTT GCT GTT GAC GAG AAT CGG GGA TCA TCG ACA CAT CAA TCA       100
Gly Ser Leu Ala Val Asp Glu Asn Arg Gly Ser Ser Thr His Gln Ser
                    15                  20                  25
TCA ACG AAA ATA TGC AGG GTG TGT GGG GAT AAG ATC GGG CAA AAG GAA       148
Ser Thr Lys Ile Cys Arg Val Cys Gly Asp Lys Ile Gly Gln Lys Glu
                30                  35                  40
AAC GGA CAA CCG TTC GTG GCT TGT CAT GTC TGT GCT TTC CCG GTT TGC       196
Asn Gly Gln Pro Phe Val Ala Cys His Val Cys Ala Phe Pro Val Cys
            45                  50                  55
CGT CCT TGT TAT GAA TAT GAA AGG AGT GAA GGA AAC CAG TGC TGT CCT       244
Arg Pro Cys Tyr Glu Tyr Glu Arg Ser Glu Gly Asn Gln Cys Cys Pro
        60                  65                  70
CAG TGC AAT ACT CGC TAT AAG CGT CAC AAA GGT AGT CCA AGA ATT TCA       292
Gln Cys Asn Thr Arg Tyr Lys Arg His Lys Gly Ser Pro Arg Ile Ser
    75                  80                  85                  90
GGA GAT GAA GAA GAT GAT TCA GAT CAA GAT GAT TTT GAT GAT GAA TTT       340
Gly Asp Glu Glu Asp Asp Ser Asp Gln Asp Asp Phe Asp Asp Glu Phe
                95                  100                 105
CAG ATT AAG AAC CGC AAG GAT GAC TCC CAT CCA CAA CAT GAA AAT GAG       388
Gln Ile Lys Asn Arg Lys Asp Asp Ser His Pro Gln His Glu Asn Glu
            110                 115                 120
GAA TAT AAT AAT AAT AAT CAT CAA TGG CAT CCC AAT GGT CAA GCT TTC       436
Glu Tyr Asn Asn Asn Asn His Gln Trp His Pro Asn Gly Gln Ala Phe
        125                 130                 135
TCA GTT GCC GGA AGC ACG GCG GGG AAG GAT TTG GAA GGG GAT AAA GAG       484
Ser Val Ala Gly Ser Thr Ala Gly Lys Asp Leu Glu Gly Asp Lys Glu
    140                 145                 150
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ATT TAC GGA AGC GAA GAA TGG AAA GAA AGA GTT GAG AAA TGG AAA GTC      532
Ile Tyr Gly Ser Glu Glu Trp Lys Glu Arg Val Glu Lys Trp Lys Val
155                 160                 165                 170
AGG CAA GAA AAA AGA GGT TTG GTA AGC AAC GAT AAT GGC GGA AAT GAT      580
Arg Gln Glu Lys Arg Gly Leu Val Ser Asn Asp Asn Gly Gly Asn Asp
                175                 180                 185
CCT CCT GAA GAA GAT GAT TAT CTC TTG GCT GAA GCT CGC CAG CCT CTA      628
Pro Pro Glu Glu Asp Asp Tyr Leu Leu Ala Glu Ala Arg Gln Pro Leu
            190                 195                 200
TGG CGA AAA GTG CCA ATT TCG TCA AGT CTG ATA AGC CCT TAC CGG ATA      676
Trp Arg Lys Val Pro Ile Ser Ser Ser Leu Ile Ser Pro Tyr Arg Ile
        205                 210                 215
GTC ATC GTC CTC CGA TTC TTC ATC CTC GCA TTT TTC CTC CGG TTC CGT      724
Val Ile Val Leu Arg Phe Phe Ile Leu Ala Phe Phe Leu Arg Phe Arg
    220                 225                 230
ATT CTA ACA CCC GCC TAC GAC GCT TAC CCG TTA TGG CTA ATC TCT GTC      772
Ile Leu Thr Pro Ala Tyr Asp Ala Tyr Pro Leu Trp Leu Ile Ser Val
235                 240                 245                 250
ATC TGC GAA GTT TGG TTC GCC TTC TCC TGG ATT CTC GAT CAG TTC CCT      820
Ile Cys Glu Val Trp Phe Ala Phe Ser Trp Ile Leu Asp Gln Phe Pro
                255                 260                 265
AAA TGG TTC CCT ATT ACT CGC GAA ACT TAC CTC GAT CGC CTC TCC TTG      868
Lys Trp Phe Pro Ile Thr Arg Glu Thr Tyr Leu Asp Arg Leu Ser Leu
            270                 275                 280
AGG TTC GAA CGT GAA GGA GAG CCC AAT CAA CTT GGC CCC GTC GAC GTC      916
Arg Phe Glu Arg Glu Gly Glu Pro Asn Gln Leu Gly Pro Val Asp Val
        285                 290                 295
TTC GTC AGT ACC GTT GAC CTT CTC AAG GAA CCC CCC ATC ATA ACC GCC      964
Phe Val Ser Thr Val Asp Leu Leu Lys Glu Pro Pro Ile Ile Thr Ala
    300                 305                 310
AAC GCG GTT CTA TCG ATC TTG GCC GTC GAT TAC CCG GTC GAG AAA GTG     1012
Asn Ala Val Leu Ser Ile Leu Ala Val Asp Tyr Pro Val Glu Lys Val
315                 320                 325                 330
TGT TGT TAT GTG TCG GAC GAT GGT GCT TCC ATG CTT CTT TTC GAT TCG     1060
Cys Cys Tyr Val Ser Asp Asp Gly Ala Ser Met Leu Leu Phe Asp Ser
                335                 340                 345
TTG TCT GAA ACG GCT GAG TTC GCG AGG AGA TGG GTT CCG TTT TGT AAG     1108
Leu Ser Glu Thr Ala Glu Phe Ala Arg Arg Trp Val Pro Phe Cys Lys
            350                 355                 360
AAG CAT AAT GTT GAG CCC AGG GCG CCG GAG TTT TAT TTC AAT GAG AAG     1156
Lys His Asn Val Glu Pro Arg Ala Pro Glu Phe Tyr Phe Asn Glu Lys
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                365                 370                 375
ATT GAT TAT TTG AAG GAC AAG GTC CAT CCT AGC TTT GTT AAA GAA CGG      1204
Ile Asp Tyr Leu Lys Asp Lys Val His Pro Ser Phe Val Lys Glu Arg
    380                 385                 390
AGA GCC ATG AAA AGG GAA TAT GAA GAA TTT AAA GTA AGG ATC AAT GCA      1252
Arg Ala Met Lys Arg Glu Tyr Glu Glu Phe Lys Val Arg Ile Asn Ala
395                 400                 405                 410
TTA GTA GCA AAA GCT CAG AAG AAA CCA GAA GAA GGA TGG GTG ATG CAA      1300
Leu Val Ala Lys Ala Gln Lys Lys Pro Glu Glu Gly Trp Val Met Gln
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[redacted]

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GAT GGC ACC CCA TGG CCC GGA AAT AAC ACT CGT GAT CAT CCT GGA ATG      1348
Asp Gly Thr Pro Trp Pro Gly Asn Asn Thr Arg Asp His Pro Gly Met
            430                 435                 440
ATT CAG GTC TAT CTA GGA AGT GCC GGT GCA CTC GAT GTG GAT GGC AAA      1396
Ile Gln Val Tyr Leu Gly Ser Ala Gly Ala Leu Asp Val Asp Gly Lys
        445                 450                 455
GAG CTG CCT CGA CTT GTC TAT GTT TCT CGT GAG AAA CGA CCT GGT TAT      1444
Glu Leu Pro Arg Leu Val Tyr Val Ser Arg Glu Lys Arg Pro Gly Tyr
    460                 465                 470
CAG CAC CAT AAG AAA GCC GGT GCT GAG AAT GCT CTG GTT CGA GTT TCT      1492
Gln His His Lys Lys Ala Gly Ala Glu Asn Ala Leu Val Arg Val Ser
475                 480                 485                 490
GCA GTG CTT ACT AAT GCA CCC TTC ATA TTG AAT CTG GAT TGT GAT CAT      1540
Ala Val Leu Thr Asn Ala Pro Phe Ile Leu Asn Leu Asp Cys Asp His
                495                 500                 505
TAC ATC AAC AAT AGC AAG GCC ATG AGG GAA GCG ATG TGC TTT TTA ATG      1588
Tyr Ile Asn Asn Ser Lys Ala Met Arg Glu Ala Met Cys Phe Leu Met
            510                 515                 520
GAT CCT CAG TTT GGA AAG AAG CTT TGT TAT GTT CAA TTT CCA CAG AGA      1636
Asp Pro Gln Phe Gly Lys Lys Leu Cys Tyr Val Gln Phe Pro Gln Arg
        525                 530                 535
TTT GAT GGT ATT GAT CGT CAT GAT CGA TAT GCT AAT CGA AAT GTT GTC      1684
Phe Asp Gly Ile Asp Arg His Asp Arg Tyr Ala Asn Arg Asn Val Val
    540                 545                 550
TTC TTT GAT ATC AAC ATG TTG GGA TTA GAT GGA CTT CAA GGC CCT GTA      1732
Phe Phe Asp Ile Asn Met Leu Gly Leu Asp Gly Leu Gln Gly Pro Val
555                 560                 565                 570
TAT GTA GGC ACA GGG TGT GTT TTC AAC AGG CAG GCA TTG TAT GGC TAC      1780
Tyr Val Gly Thr Gly Cys Val Phe Asn Arg Gln Ala Leu Tyr Gly Tyr
                575                 580                 585
GAT CCA CCA GTC TCT GAG AAA CGA CCA AAG ATG ACA TGT GAT TGC TGG      1828
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Asp Pro Pro Val Ser Glu Lys Arg Pro Lys Met Thr Cys Asp Cys Trp
                590                 595                 600
CCT TCT TGG TGT TGC TGT TGT TGC GGA GGT TCT AGG AAG AAA TCA AAG      1876
Pro Ser Trp Cys Cys Cys Cys Cys Gly Gly Ser Arg Lys Lys Ser Lys
        605                 610                 615
AAG AAA GGT GAA AAG AAG GGC TTA CTC GGA GGT CTT TTA TAC GGA AAA      1924
Lys Lys Gly Glu Lys Lys Gly Leu Leu Gly Gly Leu Leu Tyr Gly Lys
    620                 625                 630
AAG AAG AAG ATG ATG GGC AAA AAC TAT GTG AAA AAA GGG TCT GCA CCA      1972
Lys Lys Lys Met Met Gly Lys Asn Tyr Val Lys Lys Gly Ser Ala Pro
635                 640                 645                 650
GTC TTT GAT CTC GAA GAA ATC GAA GAA GGG CTT GAA GGA TAC GAA GAA      2020
Val Phe Asp Leu Glu Glu Ile Glu Glu Gly Leu Glu Gly Tyr Glu Glu
                655                 660                 665
TTG GAG AAA TCG ACA TTA ATG TCG CAG AAG AAT TTC GAG AAA CGA TTC      2068
Leu Glu Lys Ser Thr Leu Met Ser Gln Lys Asn Phe Glu Lys Arg Phe
                670                 675                 680
GGA CAA TCA CCG GTT TTC ATT GCC TCA ACT TTG ATG GAA AAT GGT GGC      2116
Gly Gln Ser Pro Val Phe Ile Ala Ser Thr Leu Met Glu Asn Gly Gly
        685                 690                 695
CTT CCT GAA GGA ACT AAT TCC ACA TCA CTG ATT AAA GAG GCC ATT CAC      2164
Leu Pro Glu Gly Thr Asn Ser Thr Ser Leu Ile Lys Glu Ala Ile His
    700                 705                 710
GTA ATT AGC TGT GGT TAT GAA GAA AAA ACT GAG TGG GGC AAA GAG ATC      2212
Val Ile Ser Cys Gly Tyr Glu Glu Lys Thr Glu Trp Gly Lys Glu Ile
715                 720                 725                 730
GGA TGG ATT TAT GGG TCG GTG ACG GAA GAT ATA TTA ACA GGT TTC AAG      2260
Gly Trp Ile Tyr Gly Ser Val Thr Glu Asp Ile Leu Thr Gly Phe Lys
                735                 740                 745
ATG CAT TGT AGA GGG TGG AAA TCG GTT TAT TGT GTA CCG AAA AGA CCG      2308
Met His Cys Arg Gly Trp Lys Ser Val Tyr Cys Val Pro Lys Arg Pro
                750                 755                 760
GCA TTC AAA GGG TCC GCT CCA ATC AAT CTC TCG GAT CGG TTG CAC CAA      2356
Ala Phe Lys Gly Ser Ala Pro Ile Asn Leu Ser Asp Arg Leu His Gln
        765                 770                 775
GTT TTG AGA TGG GCA CTT GGT TCT GTA GAA ATT TTC CTT AGT CGT CAC      2404
Val Leu Arg Trp Ala Leu Gly Ser Val Glu Ile Phe Leu Ser Arg His
    780                 785                 790
TGT CCA CTT TGG TAT GGT TAT GGT GGA AAA CTG AAA TGG CTC GAG AGG      2452
Cys Pro Leu Trp Tyr Gly Tyr Gly Gly Lys Leu Lys Trp Leu Glu Arg
795                 800                 805                 810
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CTT GCT TAT ATC AAC ACC ATT GTT TAC CCT TTC ACC TCG ATC CCT TTA      2500
Leu Ala Tyr Ile Asn Thr Ile Val Tyr Pro Phe Thr Ser Ile Pro Leu
                815                 820                 825
CTC GCC TAT TGT ACT ATT CCA GCT GTT TGT CTT CTC ACC GGC AAA TTC      2548
Leu Ala Tyr Cys Thr Ile Pro Ala Val Cys Leu Leu Thr Gly Lys Phe
            830                 835                 840
ATC ATT CCA ACT CTA AGC AAC CTT ACA AGT GTG TGG TTC TTG GCA CTT      2596
Ile Ile Pro Thr Leu Ser Asn Leu Thr Ser Val Trp Phe Leu Ala Leu
        845                 850                 855
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[redacted]

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Phe Leu Ser Ile Ile Ala Thr Gly Val Leu Glu Leu Arg Trp Ser Gly
    860                 865                 870
GTT AGC ATC CAA GAC TGG TGG CGC AAT GAA CAA TTC TGG GTG ATC GGA      2692
Val Ser Ile Gln Asp Trp Trp Arg Asn Glu Gln Phe Trp Val Ile Gly
875                 880                 885                 890
GGT GTC TCC GCC CAT CTT TTT GCT GTC TTC CAG GGC CTC CTC AAA GTC      2740
Gly Val Ser Ala His Leu Phe Ala Val Phe Gln Gly Leu Leu Lys Val
                895                 900                 905
CTA GCT GGA GTA GAC ACC AAC TTC ACC GTA ACA GCA AAA GCA GCA GAC      2788
Leu Ala Gly Val Asp Thr Asn Phe Thr Val Thr Ala Lys Ala Ala Asp
            910                 915                 920
GAT ACA GAA TTC GGT GAA CTT TAT CTC TTC AAA TGG ACA ACT CTC TTA      2836
Asp Thr Glu Phe Gly Glu Leu Tyr Leu Phe Lys Trp Thr Thr Leu Leu
        925                 930                 935
ATC CCT CCC ACA ACT CTG ATA ATA CTG AAC ATG GTC GGA GTC GTG GCC      2884
Ile Pro Pro Thr Thr Leu Ile Ile Leu Asn Met Val Gly Val Val Ala
    940                 945                 950
GGA GTT TCA GAC GCA ATC AAC AAC GGC TAT GGT TCA TGG GGT CCA TTG      2932
Gly Val Ser Asp Ala Ile Asn Asn Gly Tyr Gly Ser Trp Gly Pro Leu
955                 960                 965                 970
TTC GGC AAA CTG TTC TTC GCA TTC TGG GTC ATT CTT CAT CTT TAC CCA      2980
Phe Gly Lys Leu Phe Phe Ala Phe Trp Val Ile Leu His Leu Tyr Pro
                975                 980                 985
TTC CTC AAA GGT TTG ATG GGG AGA CAA AAC AGG ACG CCC ACC ATT GTT      3028
Phe Leu Lys Gly Leu Met Gly Arg Gln Asn Arg Thr Pro Thr Ile Val
            990                 995                 1000
GTG CTT TGG TCC ATA CTT TTG GCA TCG ATT TTC TCA CTG GTT TGG GTA      3076
Val Leu Trp Ser Ile Leu Leu Ala Ser Ile Phe Ser Leu Val Trp Val
        1005                1010                1015
CGG ATC GAT CCC TTC TTG CCC AAA CAA ACA GGT CCA GTT CTT AAA CAA      3124
Arg Ile Asp Pro Phe Leu Pro Lys Gln Thr Gly Pro Val Leu Lys Gln
```

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      1020                1025                1030
TGT GGC GTG GAG TGC TAAATGGTGT TTTACAAACC TTTCTTATTA TTTTATTTTC      3179
Cys Gly Val Glu Cys
1035
CCTTTTTGCC ACTACTGTTG ATTTGCTGTG ATTCTAAAAG GGATTTATCT TGTTTGTAAA    3239
AAGTCTCCTA TGATTTTGTT GGTTCAATTT AATTTCTATA TGGTAAAAAA ATATTTCTTT    3299
AAATTAACTA TA                                                        3311
```

(2) INFORMATION FOR SEQ ID NO: 4:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1039 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 4:

```
Met Ala Ser Thr Thr Met Ala Ala Gly Phe Gly Ser Leu Ala Val Asp
 1               5                  10                  15
Glu Asn Arg Gly Ser Ser Thr His Gln Ser Ser Thr Lys Ile Cys Arg
            20                  25                  30
Val Cys Gly Asp Lys Ile Gly Gln Lys Glu Asn Gly Gln Pro Phe Val
        35                  40                  45
Ala Cys His Val Cys Ala Phe Pro Val Cys Arg Pro Cys Tyr Glu Tyr
    50                  55                  60
Glu Arg Ser Glu Gly Asn Gln Cys Cys Pro Gln Cys Asn Thr Arg Tyr
65                  70                  75                  80
Lys Arg His Lys Gly Ser Pro Arg Ile Ser Gly Asp Glu Glu Asp Asp
                85                  90                  95
Ser Asp Gln Asp Asp Phe Asp Asp Glu Phe Gln Ile Lys Asn Arg Lys
            100                 105                 110
Asp Asp Ser His Pro Gln His Glu Asn Glu Glu Tyr Asn Asn Asn Asn
        115                 120                 125
His Gln Trp His Pro Asn Gly Gln Ala Phe Ser Val Ala Gly Ser Thr
    130                 135                 140
Ala Gly Lys Asp Leu Glu Gly Asp Lys Glu Ile Tyr Gly Ser Glu Glu
145                 150                 155                 160
Trp Lys Glu Arg Val Glu Lys Trp Lys Val Arg Gln Glu Lys Arg Gly
                165                 170                 175
Leu Val Ser Asn Asp Asn Gly Gly Asn Asp Pro Pro Glu Glu Asp Asp
            180                 185                 190
Tyr Leu Leu Ala Glu Ala Arg Gln Pro Leu Trp Arg Lys Val Pro Ile
        195                 200                 205
```

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Ser Ser Ser Leu Ile Ser Pro Tyr Arg Ile Val Ile Val Leu Arg Phe
    210                 215                 220
Phe Ile Leu Ala Phe Phe Leu Arg Phe Arg Ile Leu Thr Pro Ala Tyr
225                 230                 235                 240
Asp Ala Tyr Pro Leu Trp Leu Ile Ser Val Ile Cys Glu Val Trp Phe
                245                 250                 255
Ala Phe Ser Trp Ile Leu Asp Gln Phe Pro Lys Trp Phe Pro Ile Thr
            260                 265                 270
Arg Glu Thr Tyr Leu Asp Arg Leu Ser Leu Arg Phe Glu Arg Glu Gly
```

[redacted]

```
Glu Pro Asn Gln Leu Gly Pro Val Asp Val Phe Val Ser Thr Val Asp
    290                 295                 300
Leu Leu Lys Glu Pro Pro Ile Ile Thr Ala Asn Ala Val Leu Ser Ile
305                 310                 315                 320
Leu Ala Val Asp Tyr Pro Val Glu Lys Val Cys Cys Tyr Val Ser Asp
                325                 330                 335
Asp Gly Ala Ser Met Leu Leu Phe Asp Ser Leu Ser Glu Thr Ala Glu
            340                 345                 350
Phe Ala Arg Arg Trp Val Pro Phe Cys Lys Lys His Asn Val Glu Pro
        355                 360                 365
Arg Ala Pro Glu Phe Tyr Phe Asn Glu Lys Ile Asp Tyr Leu Lys Asp
    370                 375                 380
Lys Val His Pro Ser Phe Val Lys Glu Arg Arg Ala Met Lys Arg Glu
385                 390                 395                 400
Tyr Glu Glu Phe Lys Val Arg Ile Asn Ala Leu Val Ala Lys Ala Gln
                405                 410                 415
Lys Lys Pro Glu Glu Gly Trp Val Met Gln Asp Gly Thr Pro Trp Pro
            420                 425                 430
Gly Asn Asn Thr Arg Asp His Pro Gly Met Ile Gln Val Tyr Leu Gly
        435                 440                 445
Ser Ala Gly Ala Leu Asp Val Asp Gly Lys Glu Leu Pro Arg Leu Val
    450                 455                 460
Tyr Val Ser Arg Glu Lys Arg Pro Gly Tyr Gln His His Lys Lys Ala
465                 470                 475                 480
Gly Ala Glu Asn Ala Leu Val Arg Val Ser Ala Val Leu Thr Asn Ala
                485                 490                 495
Pro Phe Ile Leu Asn Leu Asp Cys Asp His Tyr Ile Asn Asn Ser Lys
            500                 505                 510
Ala Met Arg Glu Ala Met Cys Phe Leu Met Asp Pro Gln Phe Gly Lys
        515                 520                 525
Lys Leu Cys Tyr Val Gln Phe Pro Gln Arg Phe Asp Gly Ile Asp Arg
```